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FEATURES	JOURNAL	TITLE	AUTHORS	REFERENCE		ORGANISM	SOURCE	KEYWORDS	VERSION	ACCESSION	DEFINITION	LOCUS	AR399482	RESULT 1	
Location/Qualifiers	Patent: US 6620608-A 1 16-SEP-2003;		Gong, F., Yan, C., Di Francesco, V. and Beasley, E.M.		Unclassified.	Unknown.	Unknown.		AR399482.1 GI:40141523		Sequence 1 from patent US 6620608.	AR399482 2002 bp DNA linear PAT 18-DEC-20			
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I sogai, T. and Yamamoto, J.

S Isogai, T. and Yamamoto, J.

S Isogai, T. and Yamamoto, J.

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NEDO human cDNA sequencing project supported by Ministry of Economy, Trade and Industry of Japan; cDNA full insert sequencing: Research Association for Biotechnology (RAB); cDNA library construction: Helix Research Institute (HRI) (supported by Japan Key Technology Center etc.); 5'- & 3'-end one pass sequencing: RAB, HRI, and Biotechnology Center, National Institute of Technology and Evaluation; clone selection for full insert sequencing: HRI and RAB, annotation: HRI and RAB.
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AK095492.1 GI:21754757

oligo capping; fis (full insert sequence).

Homo sapiens (human)

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Query Match Best Local Similarity 92.0%; Pred. No. 1e-287; Matches 1509; Conservative 0; Mismatches 5; Indels 126; Gaps 1; Qy 134 GCTCTTTCACCATGCCTGGATCACTTCCTTTGAATGCAGAAGCTTGCTGGCCAAAAGATG 193	I:41002 n="MPGS MGFCTDR ESGNTDI GGVGAVA DRCYSVY NRDKNSI TSSVYGS SLCDLKS HRRTYAR	/organism="Hc /mol_type="mR /db_xref="tax /tissue_type= 221584 /codon_start= /product="3-k	purification, and characterization of recombinant purification, and characterization of recombinant of Cys129 mutant enzymes em. Biophys. 312 (1), 1-13 (1994) arce text: Homo sapiens fetal adrenal cDNA to mRNA. cation/Qualifiers	125798.1 G1:410027 3-hydroxy-3-methylglutaryl coenzyme A synthase. 3-hydroxy-3-methylglutaryl coenzyme A synthase. Homo sapiens Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eute Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Hom 1 (bases 1 to 1650) 1 (bases 1 to 1650) Rokosz,L.L., Boulton,D.A., Butkiewicz,E.A., Sanyal,G., C Lachance,P.A. and Hermes,J.D. Gran Gyforlasmic 3-bydroxy-3-methylglutaryl coenzyme A	M3H3M no sapier nplete co	65 AAAAAAAAAAAAAAAAAAAAAAAAAAAA 1994 	
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S Kalnine, N., Chen, X., Rolfs, A., Halleck, A., Hines, L., Eisenstein, S., Koundinya, M., Raphael, J., Moreira, D., Kelley, T., LaBaer, J., Lin, Y., Phelan, M. and Farmer, A.
Direct Submission

L Submitted (13-MAY-2003) BD Biosciences Clontech, 1020 East Meadow Circle, Palo Alto, CA 94303, USA
This CDS clone is a part of a collection of human full length expression clones generated by BD Biosciences Clontech and the Harvard Institute of Proteomics. Each CDS has been cloned in two forms: with and without stop-codon (to allow fusion with C-terminal tag). The CDS has been directionally cloned using BD In-Fusion(TM) cloning system between the Sall and HindIII sites of the pDNR-DUAL vector. Additional sequences in the clone: 'ACC' after Sall site and before 'ATG' to provide Kozak consensus sequence; 'GG' after last codon and before HindIII site to maintain reading frame.
Clone distribution: http://bioinfo.clontech.com/orfclones.
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Mammalia; Eutheria; Primates; Catarr
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Kalnine,N., Chen,X., Rolfs,A., Halle
Koundinya,M., Raphael,J., Moreira,D.
Phelan,M. and Farmer,A.
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                     GCTGTATATGCCACAGGAAATGCTAGACCTACAGGTGGAGTTGGAGCAGTAGCTCTGCTA
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/db_xref="GI:30583443"

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SVKTNLMQLFEESGNTDIEGIDTTNACYGGTAAVFNAVNWIESSSWDGRYALVVAGDI

AVYATGNARPTGGVGAVALLIGPNAPLIFERGLRGTHMQHAYDFYKPDMLSEYPIVDG

KLSIQCYLSALDRCYSVYCKKIHAQWQKEGNDKDFTLNDFGFMIFHSPYCKLVQKSLA

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                                      HSCOAS
H.sapiens mRNA for HMG-
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X66435.1 GI:30008
Hydroxymethylglutaryl (
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Homo sapiens
Eukaryota; Metazoa; C
Mammalia; Eutheria; P
1 (bases 1 to 1685)
Russ, A.P., Ruzicka, V.
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Catarrhini;
  Appelhans, H.
                           Vertebrata;
                   Hominidae;
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   and
   Gross, W
                            Euteleo
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t Local Similarity
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E 93041939
D 1358203
E 2 (bases 1 to 1679)
S Russ,A.
Direct Submission
L Submitted (26-MAY-1992) A. Russ, Labor fur angewandte Biochemie, Theodor Stern-Kai 7, W-6000 Frankfurt am Main 70, FRG revised by [2]
E 3 (bases 1 to 1685)
S Russ,A.
Direct Submission
L Submitted (10-AUG-1992) Andreas Russ, Zentrum der biologischen Chemie, J.W.-Goethe-Universitaet Frankfurt, Theodor-Stern-Kai 7, Frankfurt, 6000, Germany Location/Qualifiers
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50	448 TGGATGAAGGAGTAGGACTTGTGCATTCAAACATAGCAACTGAGCATATTCCAAGCCCT
1447 1551	1388 GGGTGGATGAAAAGCACAGAAGAACTTACGCTCGGCGTCCCACTCCAAATGATGACACTT
1387 1491	1328 TCAACTATATTCCCCAGGGTTCAATAGATTCACTCTTTGAAGGAACGTGGTACTTAGTTA
1327 1431	1268 GTGTGGCACCAGATGTCTTCGCTGAAAACATGAAGCTCAGAGAGGACACCCATCATTTGG
1267 1371	1208 CTCTTGATAAATAACAGCAAGTTTATGTGATCTTAAATCAAGGCTTGATTCAAGAACTG
1207 1311	1148 CTGGTTTGGCTGCCACTCTGTACTCTCTTAAAGTCACACAAGATGCTACACCGGGGTCTG
1147 1251	1088 TAGCACAGTACTCACCTCAGCAATTAGCAGGGAAGAGAATTGGAGTGTTTTCTTATGGTT
1087 1191	1028 TATCAAATCAAAATGGAAATATGTACACATCTTCAGTATATGGTTCCCTTGCATCTGTTC
1027 1131	968 AGGCATTTATGAAGGCTAGCTCTGAACTCTTCAGTCAGAAAACAAAGGCATCTTTACTTG
967 1071	908 GCCTGGAAGCCTTTGGGGATGTTAAATTAGAAGACACCTACTTTGATAGAGATGTGGAGA
907	848 GGATGTTGCTGAATGACTTCCTTAATGACCAGAATAGAGATAAAAATAGTATCTATAGTG
847 951	788 ATTTTGGCTTCATGATCTTTCACTCACCATATTGTAAACTGGTTCAGAAATCTCTAGCTC
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L Proc. Natl. Acad. Sci. U....

E 2388257

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E 2 (bases 1 to 3260)

SS Strausberg,R.

Direct Submission

AL Submitted (06-MAY-2002) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Cancer Genomics Offi
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S Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
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Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,
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Fahey, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S.,
Sanchez, A., Whiting, M., Madan, A., Woung, A.C., Shevchenko, Y.,
Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, B.D.,
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Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E.,
Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.
Generation and initial analysis of more than 15,000 full-length
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                                                                                                                                                                                                                   Contact: MGC help desk
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLM
DNA Sequencing by: Baylor College of Medicine Human Ger
Sequencing Center
Center code: BCM-HGSC
Web site: http://www.hgsc.bcm.tmc.edu/cdna/
Contact: amg@bcm.tmc.edu
Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Lou
Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., Na
A.N., Gibbs, R.A.
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Mammalia; E
Clone distribution: MGC clone distribution information can through the I.M.A.G.E. Consortium/LLNL at: http://image.lli Series: IRAK Plate: 53 Row: p Column: 14
This clone was selected for full length sequencing because passed the following selection criteria: matched mRNA gi:
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                                                                                                              ion information can be for at: http://image.llnl.gov
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GAGACAATCATCGACAAATCGAAATCAGTGAAGTCTAATTTGATGCAGCTGTTTGAGGÁG 420 TCTGGGAATACAGATATAGAAGGAATCGACAACCAACTAATGCATGC	· 44	AATCTTATGGAGAGAAATAACCTTTCCTATGATTGCATTGGGCGGCTGGAAGTTGGAACA 426 	AAGATGGGCTTCTGCACAGATAGAGAAGATATTAACTCTCTTTGCATGACTGTGGTTCAG 366 	GAGTTGGAAAAATATGATGGTGTAGATGCTGGAAAGTATACCATTGGCTTGGGCCAGGCC 306 	AAAGATGTGGGAATTGTTGCCCTTGAGATCTATTTTCCTTCTCAATATGTTGATCAAG 	CACGCTTGCTCTTTCACCATGCCTGGATCACTTCCTTTGAATGCAGAAGCTTGCTGGCCA 186	GGGGAGGCGCCGCGACTGTCCTTTCGTGGCTCACTCCCTTTCCTCTGCTGCCGC	Simi 2;	/note="HMG_CoA_synt; Region: Hydroxymethylglutaryl-coenzyme A synthase" /db_xref="CDD:pfam01154"	TPGSALDKITASLCDLKSRLDSRTCVAPDVFAENMKLREDTHHLANYIPQCSIDSLFE GTWYLVRVDEKHRRTYARRPFTNDHSLDEGMGLVHSNTATEHIPSPAKKVPRLPATSA ESESAVISNGEH" ature 1151485	ALYATGNAKETGGVGAVALLIGENAPLIEDRGLRGTHMQHAYDFYKPDMLSEYPVVDG KLSIQCYLSALDRCYSVYRKKIRAQWQKEGKDKDFTLNDFGFMIFHSPYCKLVQKSLA RMFLNDFLNDQNRDKNSIYSGLEAFGDVKLEDTYFDRDVEKAFMKASSELFNQKTKAS LLVSNQNGNMYTSSVYGSLASVLAQYSPQQLAGKRVGVFSYGSGLAATLYSLKVTQDA	/ db_xrer="Locusid: 208715" /translation="MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDA /translation="MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDA GKYTIGLGQARMGFCTDREDINSLCLTVVQKLMERHSLSYDCIGRLEVGTETIIDKSK SVKSNLMQLFEESGNTDIEGIDTTNACYGGTAAVFNAVNWVESSSWDGRYALVVAGDI	/codon_scarc=1 /product="Hmgcs1 protein" /protein_id="AAH29693.1" /db_xref="GI:20988709"	/db_xref="MGI:107592" 791641	e="Hmgcsl" e="synonym: MGC	3260	mouse. Taken by biopsy." ne_lib="NCI_CGAP_Mam2"	ne="MGC:36525 IMAGE:5375374" sue_type="Mammary tumor. MMTV-LTR/INT3 model	type="mRNA" type="mRNA" ain="FVB/N-3"	ation/Qualis
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THORS Strausberg, R. L., Feingold, E. A., Grouse, L. H., Derge, J. G., Klausberg, R. L., Feingold, E. A., Grouse, L. H., Derge, J. G., Altschul, S. F., Zeeberg, B., Buetow, K. H., Schaefer, C. F., Batt, N. Hopkins, R. F., Jordan, H., Moore, T., Max, S. I., Wang, J., Hsieh, F. Stapleton, M., Soares, M. B., Bonaldo, M. F., Casavant, T. L., Stepleton, M., Soares, M. B., Bonaldo, M. F., Casavant, T. L., Scheetz, T. E., Brownstein, M. J., Usdin, T. B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S. S., Loquellano, N. A., Peters, G. J. Abramson, R. D., Mullahy, S. J., Bosak, S. A., McEwan, P. J., Worley, K. C., Hale, S., Garcia, A. M., Gay, L. J., Hulyk, S. M., Villalon, D. K., Muzny, D. M., Sodergren, E. J., Lu, X., Gibbs, R. A., Fahey, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S., Sanchez, A., Whiting, M., Madan, A., Young, A. C., Shevchenko, Y., Buffard, G. G., Blakesley, R. M., Touchman, J. W., Green, E. D., Dickson, M. C., Rodriguez, A. C., Grimwood, J., Schmutz, J., Myers, R. M. Schmerch, A., Schein, J. E., Jones, S. J. and Marra, M. A. Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences

Proc. Matl. Acad. Sci. U.S. A. 99 (26), 16899-16903 (2002)

Strausberg, R. Direct s. P. Direct s. P. C. Shevel L. C. Shevel 
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NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: The Cepko Laboratory
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Gunaratne, P.H., Garcia, A.M., Lu, X., Hul: Kowis, C.R., Sneed, A.J., Martin, R.G., Mu A.N., Gibbs, R.A.
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Conservative
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/clone="MGC:36662 IMAGE:5366786"

/tissue_type="Eye, retina, mouse s

/clone_lib="NIH_MGC_94"

/lab_host="DH10B"

/note="Vector
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/db_xref="LocusID:208715"
/db_xref="MGI:107592"
25._.1587
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Hydroxymethylglutaryl-coenzyme
/db_xref="CDD:pfam01154"
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1206	TAIGGITCIGGITIGGCIGCCACICIGIACICITIAAAGICACACAAGAIGCIACACCACACACACACACACAC	114	β ξ
14	TCTGTCCTGGCACAGTACTCACCTCAGCAGTTGGCAGGGAAGAGGGGTTGGAGTGTTCTCT	80	В
4	GCAGGGAAGAGAATTGGAGTGTTTTCT	1081	γ <u>ο</u>
1086	7 TIGCTIGTGTCTAATCAGAATGGAAATATGTACACATCCTCTGTCTATGGTTCCCTGGCT	1027	DЬ
1080	TACTTGTATCAAATCAAAATGGAAATATGTACACATCTTCAGTATATGGTTCCCTTGCA	102:	Ş
1026	CATTTATGAAGGCTAGTTCTGAGCTATTCAACCAGAAAACAAAGGCGTCT	967	da
1020	GGAGAAGGCATTTATGAAGGCTAGCTCTGAACTCTTCAGTCAG	961	γQ
966		907	dd
960	ATAGTGGCCTGGAAGCCTTTGGGGATGTTAAATTAGAAGACACCTACTTTGATAGAGAT	901	Ş
906		847	망
900	TAGCTCGGATGTTGCTGAATGACTTCCTTAATGACCAGAATAGAGATAAAAAATAGTATC	841	γQ
846	GGTGCAGAAATCT	787	dd
840	AATGATTTTGGCTTCATGATCTTTCACTCACCATATTGTAAACTGGTTCAGAAATCT	783	Ş
786	7 GTCTACCGCAAAAGATCCGTGCCCAGTGGCAGAAAGAGGGAAAGGATAAAGATTTTACC	72.	Дb
780	TCTACTGCAAAAAGATCCATGCCCAGTGGCAGAAAGAGGGGAAATGATAAAGATTTTA	72:	γQ
726	7 GTGGTCGACGGAAAGCTCTCCATACAGTGCTACCTCAGCGCCCTGGACCGCTGCTATTCT	66.	Дb
720	TAGATGGAAAACTCTCCATACAGTGCTACCTCAGTGCATTAGACCGCTGCTACTCT	661	γQ
666	CACACATGCAGCATGCCTATGACTTTTACAAGCCTGACATGCTCTCCGAGTACCCT	607	дb
660	GGACACATATGCAACATGCCTATGATTTTTACAAGCCTGATATGCTATCTGAATATCCT	601	γQ
606	TGTGGCCCTGCTAATTGGGCCAAACGCTCCTCTAATTTTTGACCGAGGGCTCCGT	547	Дb
600	CTTCGT	592	γΩ
546	TGCAGGAGACATTGCTATATATGCCACAGGAAATGCCAGACCTACAGGTGGAGTT	487	DЬ
591		592	γQ
486	GGACGATATGCTCTG	427	ф
591	CTGTCTTCAATGCTGTTAACTGGATTGAGTCCAGCTCTTGGGAT	547	Ş
426		367	40
546	CTGGGAATACAGATATAGAAGGAATCGACACAACTAATGCATGC	487	δ
366		307	Дb
486	ACAATCATCGACAAATCAAAGTCTGTGAAGACTAATTTGATGCAGCTGTTTGAAGAG	427	Ş
306		247	ф
426	CTTATGGAGAGAATAACCTTTCCTATGATTGCATTGGGCGGCTGGAAGTTGGAACA	367	Ş
246	GATCGTGAAGACATCAACTCTCTTTGCCTGACTGTGGTTCAG	187	Дb
366	, АТСССТТСТССАСАСАТАСАВАТАТТААСТСТСТТТССАТСАСТСТСТСТ	307	γQ
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306	agttggaaaaatatgatggtgtagatgctggaaagtataccattggcttgggccaggcc	247	γQ

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ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM REFERENCE AUTHORS

RESULT 9
BC023851
LOCUS
DEFINITION

BC023851

BC023851

ON Mus musculus 3-hydroxy-3-methylglutaryl-Coenzyme A synthase 1, mRNA (cDNA clone MGC:36620 IMAGE:5347038), complete cds.

BC023851

BC023851

BC023851.1 GI:23271493

MGC.

Mus musculus (house mouse)

Mus musculus (house mouse)

Mus musculus (house mouse)

Mus musculus (house mouse)

SM Mus musculus (house mouse)

Mus musculus (house mouse)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

E 1 (bases 1 to 2703)

S Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,

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PUBMED
REFERENCE
AUTHORS
TITLE
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COMMENT
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Strausberg, R.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc_mgc@nhgri.nih.gov
Akhter,N., Ayele,K., Beckstrom-Sternberg,S.M., Benjamin,B.,
Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C., Brooks,S.,
Dietrich,N.L., Granite,S., Guan,X., Gupta,J., Haghighi,P.,
Hansen,N., Ho,S.-L., Karlins,E., Kwong,P., Laric,P., Legaspi,R.,
Maduro,Q.L., Masiello,C., Maskeri,B., Mastrian,S.D.,McCloskey,J.C.,
McDowell,J., Pearson,R., Stantripop,S., Thomas,P.J., Touchman,J.W.,
Tsurgeon,C., Vogt,J.L., Walker,M.A., Wetherby,K.D., Wiggins,L.,
Young,A., Zhang,L.-H. and Green,E.D.
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/db_xref="LocusID:208715
/db_xref="MGI:107592"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               organism="Mus musculus
/mol_type="mRNA"
/strain="FVB/N"
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                                                                                                                                                                                                                                                                                             gene="Hmgcs1"
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'lab_host="DH10B"
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protein_id="AAH23851.1"
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db_xref="LocusID:208715"
translation="MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDA
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                                     GCAAAAAGATCCATGCCCAGTGGCAGAAAGAGAGGGAAATGATAAAGATTTTACCTTGAATG
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SVKSNLMQLFEESGNTDIEGIDTTNACYGGTAAVFNAVNWVESSSWDGRYALVVAGDI
AIYATGNARPTGGVGAVALLIGPNAPLIFDRGLRGTHMQHAYDFYKPDMLSEYPVVDG
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ESESAVISNGEH"
185. .1555
/note="HMG_CoA_synt; Region:
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/db_xref="CDD:pfam01154"
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Mus musculius (house mouse)

Mus musculius (house mouse)

Mus musculius (Chordata, Craniata, Vertebrata, Euteleostomi, Mammalia, Butheria, Rodentia, Sciurognathi, Muridae, Murinae, Mus.

B 1 (bases 1 to 378)

Strausberg, R.D., Collins, F.S., Wagner, L., Shemen, C.M., Schuler, G.D., Altschin, S.F., Joedbarg, B., Buecow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.D., Collins, F.S., Wagner, L., Shemen, C.M., Schuler, G.D., Altschin, S.F., Joedbarg, B., Buecow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.D., Mollins, R.D., Martin, J., Usdan, T.B., Toshiyuki, S., Strapleton, M., Soares, M.B., Bonaddo, M.F., Casavant, T.L., Scheefer, T.B., Brownsteen, M.J., Usdan, T.B., Toshiyuki, S., Carninci, P., Frange, C., Raha, S.S., Loquellano, N.A., Peters, G.J., Abrameon, R.D., Mullahy, S.J., Loquellano, N.A., Peters, G.J., Abrameon, R.D., Millahy, S.J., Loquellano, N.A., Peters, G.J., McKernan, K.J., Malek, J.A., Gunarathe, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalon, D.K., Muzny, D.M., Sodergren, B.J., Lu, X., Gibbs, R.A., Villalon, D.K., Muzny, D.M., Sodergren, B.J., Lu, X., Gibbs, R.A., Villalon, D.K., Muzny, D.M., Sodergren, B.J., Lu, X., Gibbs, R.A., Villalon, D.K., Muzny, D.M., Sodergren, B.J., Lu, X., Green, B.D., Santhaz, A., Willalon, D.K., Muzny, D.M., Sodergren, S.J., Lu, X., Green, B.D., D.S., Sanchaz, A., Willalon, D.K., Muzny, D.M., Sodergren, S.J., Lu, X., Shwitento, Y., Shwite, J., Schmutz, J., Wyers, R.M., Staleka, W., Staleka, W., Staleka, W., Staleka, W., Staleka, W., Schmutz, J., Wyers, R.M., Staleka, W., Schmutz, J., Wyers, R.M., Butterfield, Y.S., Kzzyvinski, M.J., Skaleka, W., Schmutz, J., Wyers, R.M., Butterfield, Y.S., Kzzyvinski, M.J., Skaleka, W., Schmutz, J., Wyers, R.M., Butterfield, Y.S., Kzzyvinski, M.J., Skaleka, W., Schmutz, J., Wyers, R.M., Butterfield, Y.S., Kzzyvinski, M.J., Skaleka, W., Schwatz, J., Wyers, R.M., Butterfield, Y.S., Kzzyvinski, M.J., Skaleka, W., Skaleka, W., Skaleka, W., Skaleka, W., Skaleka, W.
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(cDNA clone M
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BC031363.1 G
MGC.
                                                                   Clone distribution: MGC clone distribution information can be for through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Series: IRAK Plate: 54 Row: e Column: 15
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 221223.
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/organism="Mus musculus"
/mol_type="mRNA"
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RMFLNDFLNDQNRDKNSIYSGLEAFGDVKLEDTYFDRDVEKAFMKASSELFNQKTKAS
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TPGSALDKITASLCDLKSRLDSRTCVAPDVFAENMKLREDTHHLANYIPQCSIDSLFE
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117. .1487
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Hydroxymethylglutaryl-coenzyme A synthase"
/db_xref="CDD:pfam01154"
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 GCTTCTGCACAGATAGAGAAGATATTAACTCTCTTTGCATGACTGTGGTTCAGAATCTTA
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Rattus norvegicus (Norway rat)

M Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae
Rattus.

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Ayte,J., Gil-Gomez,G. and Hegardt,F.G.
Nucleotide sequence of a rat liver cDNA encoding th
3-hydroxy-3-methylglutaryl coenzyme A synthase
Nucleic Acids Res. 18 (12), 3642 (1990)
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(b 1-38,40,41-62,64-3275)
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Pirect Submission
Submitted (17-APR-1990) Hegardt F.G., University of Barcelona, 1 of Biochemistry, School of Pharmacy, Placa Pius XII, s/n. 08028 barcelona, Spain
(b 1-38,40,41-62,64-3275)
3 (bases 1 to 3275)
Haegardt, F.G.
Direct Submission
Submitted (30-JUL-1990)
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RESULT 14 AL356361 AL356361 AL356361 AL356361 AL356361 DEFINITION Human DNA sequence from clone RP11-35L17 on chromosome 1, complete sequence. ACCESSION AL356361 VERSION KEYWORDS SOURCE ORGANISM HTG. SOURCE ORGANISM Homo sapiens (human) ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 127832) Lloyd,C. TITLE JOURNAL Cambridgeshire, CB10 1SA, UK. B-mail enquiries: humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk On Jun 21, 2002 this sequence version replaced gi:18693024. Unity sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the	Db 1341 GTGTGGCACCAGACGTCTTTGCAAAACATGAAGGTCAGAGAGGACACACAC

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This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/Chr1

RP11-35L17 is from the library RPCI-11.1 constructed by the group of Pieter de Jong. For further details see http://www.chori.org/bacpac/home.htm

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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-35L17"
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Birren,B., Linton,L., Nusbaum
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O sapiens (human)
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                            Center clone name: 661 L 5
----- Summary Statistics
Sequencing vector: M13; M77815;
Sequencing vector: Plasmid; n/a
0.524475524475524Chemistry: Dye
                                                                                         Center: Whitehead Institute/ MIT Center code: WIBR Web site: http://www-seq.wi.mit.
                                                                   Contact: sequence_submissions@genome.wi.mit ----- Project Information
Center project name: L4319
Assembly program:
Consensus quality:
Consensus quality:
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REFERENCE AUTHORS TITLE JOURNAL REFERENCE AUTHORS

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ACCESSION VERSION KEYWORDS

1199 CGGGGTCTGCTCTTGATAAAATAACAGCAAGTTTATGTGATCTTAAATCAAGGCTTGATT 1258 	30	GGAAAAA TATGATGGTGTAGATGCTGGAAAGTATACCATTGGCTTGG(245
1139 CTTATGGTTCTGGTTTGGCCGCCACTCTGTACTCTCTTAAAGTCACACAAGATGCTACAC 1198	gaps 6:	55.1%; Score 1103.2; DB 2; Length 170834; arity 79.4%; Pred. No. 1.2e-229; onservative 0; Mismatches 198; Indels 183; G	Query Match Best Local Simila Matches 1465; Co
CCCAGCGCAGTACTCACCTCAGCAATGATCAGGGAAGAG-ACTGGAGTC	מֹם	note="assembl	
1079 CATCTGTTCTAGCACAGTACTCACCTCAGCAATTAGCAGGGAAGAGAGAATTGGAGTGTTTT 1138	Qy	41035153563 note="assembly_fra 53664170834	misc_reature
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Run on: OM protein 1 protein search, June 24, 2004, 12:56:26 Copyright GenCore (c) 1993 using sw model version - 2004 ; Search time 44 Seconds {without alignments} 565.671 Million cell updates/sec 5.1.6 Compugen Ltd

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Scoring table: BLOSUM62 Gapop 10.0 , Gapext 0.5

Total number Searched: of hits satisfying chosen parameters: 141681 seqs, 52070155 residues

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Minimum Maximum BQ BQ seq length:
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Post-processing: Minimum Match 0% Maximum Match 100% Listing first 45 summaries

Database SwissProt_42:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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EMBL; BC000297; AAH00297.1; -.

PIR; S27197; S27197.

PIR; S45497; S45497.

Genew; HGNC:5007; HMGCS1.

MIM; 142940; -.

GO; GO:0005737; C:cytoplasm; TAS.

GO; GO:0004421; F:hydroxymethylglutaryl-ConterPro; IPR000590; HMG_COA_synt_AS.

InterPro; IPR008260; HMG_COA_synt_AS.

InterPro; IPR008260; HMG_COA_synthASE; 1.

PROSITE; PS01126; HMG_COA_synthesis; Mulacyl-Conflict

ACT_SITE 129 129

MUTAGEN 129 129

CONFLICT 248 248 G -> A (IN RICONFLICT 251 251 K -> V (IN RICONFLICT 364 364 Q -> H (IN RICONFLICT 420 420 P -> Q (IN RICONFLICT 519 520 EH -> VW (IN RICONFLICT 519 520 EH
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RESTRAIN-Sprague Dawley; TISSUE=Liver;

RA MEDLINE=90301491; pubMed=1972979;

RA MEDLINE=90301491; pubMed=1972979;

RA Ayte J., Gil-Gomez G., Hegardt F.G.;

RI "Nuclectide sequence of a rat liver CDNA encoding the cytosolic 3-

RT "Nuclectide sequence of a rat liver CDNA encoding the cytosolic 3-

RT Nucleic Acids Res. 18:3642-3642(1990).

CC -!- FUNCTION: This enzyme condenses acetyl-COA with acetoacetyl-COA to

form HMG-COA, which is the substrate for HMG-COA reductase.

CC -!- FUNCTION: This enzyme condenses acetyl-COA acetoacetyl-COA = (S)-3-

hydroxy-3-methylglutaryl-COA + (COA) + ACETOACETYL-COA = (S)-3-

hydroxy-3-methylglutaryl-COA + COA.

CC -!- FUNCTION: CACTIVITY: ACETYL-COA + H(2)O + acetoacetyl-COA = (S)-3-

hydroxy-3-methylglutaryl-COA + COA.

CC -!- FUNCTION: Cytoplasmic.

CC -!- SIMILARITY: Belongs to the HMG-COA PRIOR TO THE

CC -!- SIMILARITY: Belongs to the HMG-COA synthase family.

CC -!- SIMILARITY: Belongs to the HMG-COA synthase family.

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01-AUG-1990 (Rel. 15, Last sequence update)
10-OCT-2003 (Rel. 42, Last annotation update)
Hydroxymethylglutaryl-CoA synthase, cytoplasmic (Esynthase) (3-hydroxy-3-methylglutaryl coenzyme A seynthase) (Ratus norvegicus (Rat).
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata Mammalia; Eutheria; Rodentia; Sciurognathi; Murida NCBI_TaxID=10116;
                                                                                                                                                                                                                                                                                                                            EMBL; X52625; CAA36852.1; -.

PIR; S12736; S12736.

InterPro; IPR000590; HMG_COA_synt AS.

InterPro; IPR008260; HMG_COA_synth.

Pfam; PF01154; HMG_COA_synt; 1.

PROSITE; PS01226; HMG_COA_SYNTHASE; 1.

Transferase; Cholesterol biosynthesis; Multigene ACT_SITE 129 POTENTIAL.

SEQUENCE 520 AA; 57433 MW; CB213A27B0C177CB C
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Theory (Rel. 13, Created)

Tol-JAN-1990 (Rel. 13, Last sequence update)

Tol-JAN-1990 (Rel. 13, Last sequence update)

Tol-JAN-1990 (Rel. 14, Last sequence update)

Tol-OCT-2003 (Rel. 14, Last sequence update)

Expithase) (3-hydroxy-3-methylglutaryl coenzyme A synthase).

MREDINIBE-OVATY;

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EMBL; L00331; AAA37076.1; JOINED.
EMBL; L00332; AAA37076.1; JOINED.
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EMBL; L00333; AAA37076.1; JOINED.
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ENBL; L00333; AAA37076.1; JOINED.
ENBL; L00333; AAA37076.1; JOINED.
ENBL; L00332; AAA37076.1; JOINED.
ENBL; 
01-NOV-1991 (Rel. 20, Created)
01-NOV-1991 (Rel. 20, Last sequence update)
10-OCT-2003 (Rel. 42, Last annotation update)
1ydroxymethylglutaryl-CoA synthase, cytoplasmic (Esynthase) (3-hydroxy-3-methylglutaryl coenzyme A synthase) (3-hydroxy-3-methylglutaryl coenzyme A synthase)
1MGCS1 OR HMGCS.
3allus gallus (Chicken).
3allus gallus (Chicken).
4rchosauria; Aves; Neognathae; Galliformes; Phasia
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InterPro; IPR000590; HMG_CoA_synth.

InterPro; IPR008260; HMG_CoA_synth.

Pfam; PF01154; HMG_CoA_synt; 1.

PROSITE; PS01226; HMG_COA_SYNTHASE; 1.

PROSITE; PS01226; HMG_COA_SYNTHASE; 1.

Transferase; Cholesterol biosynthesis; Multigene far POTENTIAL.

Transferase; Cholesterol biosynthesis; Multigene far POTENTIAL.
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                                          PDVFAENMKLREDTHHLVNYIPQGSIDSLFEGTWYLVRVDEKHRRTYARRPTPNDDTLDE
                                                                           YSPEHLAGQRISEFSYGSGFAATLYSIRVTQDATPGSALDKITASLSDLKARLDSRKCIA
                                                                                          EGIDTTNACYGGTAAVFNAVNWIESSSWD
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Pred. No. 2e-1
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RESIDUE-Brain;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
Strausberg R.L., Feingold E.A., Grouse L.H., Shenmen C.M., Schuler G.D.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Altschul S.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Ketteman M., Madan A., Rodrigues S., Sanchez A.,
RA Hotriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
"Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
                                                             cloning, chromosome warrant vertebrate evolution.";
vertebrate evolution.";
Genomics 23:552-559(1994).
-!- FUNCTION: This enzyme condenses acetyl-CoA with acetoacetyl-CoA to form HMG-CoA, which is the substrate for HMG-CoA reductase.
-!- CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acetoacetyl-CoA = (S)-3-hydroxy-3-methylglutaryl-CoA + CoA.
-!- PATHWAY: RESPONSIBLE, TOGETHER WITH HMG-COA LYASE FOR KETONE
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01-OCT-1996 (Rel. 34, Last sequence update)
10-OCT-2003 (Rel. 42, Last annotation update)
Hydroxymethylglutaryl-CoA synthase, mitochondrial precursor
(EC 2.3.3.10) (HMG-CoA synthase) (3-hydroxy-3-methylglutaryl coenzyme A synthase).
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MEDLINE=95200282; PubMed=7893153;
Mascaro C., Buesa C., Ortiz J.A., Haro D., Hegardt
Mascaro C., Buesa C., Ortiz J.A., Haro D., Hegardt
"Molecular cloning and tissue expression of human r
hydroxy-3-methylglutaryl-CoA synthase.";
Arch. Biochem. Biophys. 317:385-390(1995).
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Boukaftane Y., Mitchell G.A.;
"Cloning and characterization of the human mitochondrial 3-hydroxy-3-methylglutaryl CoA synthase gene.";
Gene 195:121-126(1997).
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Mammalia; Eutheria; Primates;
NCBI_TaxID=9606;
                                                                                                                                                                                           MEDLINE=95154824; PubMed=7851882;
Boukaftane Y., Duncan A., Wang S., Labuda D.,
Sarrazin J., Schappert K., Mitchell G.A.;
"Human mitochondrial HMG CoA synthase: liver
cloning, chromosome mapping to 1p12-p13, and
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is SWISS-PROT entry is copyright. It is produced through a collaboration tween the Swiss Institute of Bioinformatics and the EMBL outstation - European Bioinformatics Institute. There are no restrictions on its by non-profit institutions as long as its content is in no way lifted and this statement is not removed. Usage by and for commercial ities requires a license agreement (See http://www.isb-sib.ch/announce/send an email to license@isb-sib.ch).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  e J., Gil-Gomez G., Haro D., Marrero P.F., Hegardt F.G.; t mitochondrial and cytosolic 3-hydroxy-3-methylglutaryl-CoA thases are encoded by two different genes."; c. Natl. Acad. Sci. U.S.A. 87:3874-3878(1990). FUNCTION: This enzyme condenses acetyl-CoA with acetoacetyl-CoA form HMG-CoA, which is the substrate for HMG-CoA reductase. CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acetoacetyl-CoA hydroxy-3-methylglutaryl-CoA + CoA.
PATHWAY: RESPONSIBLE, TOGETHER WITH HMG-COA LYASE FOR KETONE
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; A35865; A35865.
erPro; IPR000590; HMG_COA_synt AS.
erPro; IPR008260; HMG_COA_synth.
m; PF01154; HMG_COA_synt; 1.
SITE; PS01226; HMG_COA_SYNTHASE; 1.
nsferase; Cholesterol biosynthesis; Mitochondrion;
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malia; Eutheria;
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AUG-1991 (Rel. 19, Last sequence update)
OCT-2003 (Rel. 42, Last annotation update)
roxymethylglutaryl-CoA synthase, mitochondrial precursor
2.3.3.10) (HMG-CoA synthase) (3-hydroxy-3-methylglutaryl
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LINE=90251660; PubMed=1971108;
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SUBCELLULAR LOCATION: Mitochondrial.
TISSUE SPECIFICITY: Liver and kidney.
SIMILARITY: Belongs to the HMG-CoA synthase family.
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STRAIN=CHS; TISSUE=Liver;

MEDLINE=95154824; PubMed=7851882;

MEDLINE=95154824; PubMed=7851882;

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 EMBL; U12790; AAA92675.1; -.
EMBL; U12791; AAA92676.1; -.
SWISS-2DPAGE; P54869; MOUSE.
MGD; MGI:101939; Hmgcs2.
InterPro; IPR000590; HMG_CoA_sy
Pfam; PF01154; HMG_CoA_synt; 1.
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Eukaryota; Metazoa; (
Mammalia; Eutheria; F
NCBI_TaxID=10090;
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P54869; Q64740;
01-OCT-1996 (Rel. 34, Created)
01-OCT-1996 (Rel. 34, Last sequence update)
10-OCT-2003 (Rel. 42, Last annotation update)
Hydroxymethylglutaryl-CoA synthase, mitochondrial precursor (EC 2.3.3.10) (HMG-CoA synthase) (3-hydroxy-3-methylglutaryl
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Adams S.H., Alho C.S., Asins G., Hegardt F.G., Nadams S.H., Alho C.S., Nadams G., Hegardt F.G., Nadams S.H., Alho C.S., Nadams G., Hegardt F.G., Nadams G., Hegardt F.G., Nadams G., Hegardt F.G., Nadams G., N
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Eukaryota; Metazoa; (Mammalia; Eutheria; NCBI_TaxID=9823; [1]
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HMCM_PIG STANDARD; PRT; 508 AA.

15-DEC-1998 (Rel. 37, Created)
15-DEC-1998 (Rel. 37, Last sequence update)
10-OCT-2003 (Rel. 42, Last annotation update)
Hydroxymethylglutaryl-CoA synthase, mitochondrial precursor
(EC 2.3.3.10) (HMG-CoA synthase) (3-hydroxy-3-methylglutaryl
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       ne condenses acetyl-CoA with acetoacetyl-CoA is the substrate for HMG-CoA reductase.
Acetyl-CoA + H(2)O + acetoacetyl-CoA = (S)-3
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01-OCT-1996 (Rel. 34, Last sequence upd:
10-OCT-2003 (Rel. 42, Last annotation upd:
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InterPro; IPR008260; HMG_CoA_sy
Pfam; PF01154; HMG_COA_synt; 1.
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s SWISS-PROT entry is copyright. It is produced through a collaboration: ween the Swiss Institute of Bioinformatics and the EMBL outstation - European Bioinformatics Institute. There are no restrictions on its by non-profit institutions as long as its content is in no way lified and this statement is not removed. Usage by and for commercial ities requires a license agreement (See http://www.isb-sib.ch/announce/send an email to license@isb-sib.ch).
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 L; X73679; CAA52032.1; -.
; S38986; S38986.
erPro; IPR000590; HMG_COA_synth.
m; PF01154; HMG_COA_synt; 1.
SITE; PS01226; HMG_COA_SYNTHASE; 1.
nsferase; Sterol biosynthesis; Multigene family.
SITE 116 116 POTENTIAL.
UENCE 453 AA; 50332 MW; C79EB2376270F348 CRC64;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      ttella germanica (German cockroach).
aryota; Metazoa; Arthropoda; Hexapoda; Insecta;
ptera; Orthopteroidea; Dictyoptera; Blattaria;
ttellidae; Blattellinae; Blattella.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        J. Biochem. 217:691-699(1993).

FUNCTION: This enzyme condenses acetyl-CoA with acetoacetyl-CoA form HMG-CoA, which is the substrate for HMG-CoA reductase (By similarity).

CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acetoacetyl-CoA = (S)-3 hydroxy-3-methylglutaryl-CoA + CoA.

PATHWAY: PRODUCTION OF MEVALONATE FROM HMG-COA PRIOR TO THE SYNTHESIS OF STEROLS AND ISOPRENOIDS SUCH AS JUVENILE HORMONES.

SIMILARITY: Belongs to the HMG-CoA synthase family.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           53.9%;
Il Similarity 56.6%;
259; Conservative 66
                                                                                                                                                                                                                                                                                                                                                                                                                                                                         RDVEKAFMKASSELFSOKTKASLLVSNONGNMYTSSVYGSLASVLAQYSPOOLAGKRIGV
                                                                                                                                                                                                                                                                                                                                                       TAAVFNAVNWIESSSWD---
                                                                                                                                                      3-methylglutaryl-coenzyme-A synthase Cloning, expression, developmental particles
                                                                            KAVMTYSKNMFEEKTKPS
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J., Buesa C., Piulachs
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Pred. No. 3.3e-91;
66; Mismatches 84;
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                                                                             TPSLYGGLVSLLVSKSAQELAGKRVAL
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pattern and tissue
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Blaberoidea;
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SEQUENCE FROM N.A.
MEDLINE=94216267; PubMed=7909314;
Buesa C., Martinez-Gonzalez J., Casals N., H.
Buesa C., Martinez-Gonzalez J., Casals N., H.
Belles X., Hegardt F.G.;
"Blattella germanica has two HMG-CoA synthas,
regulated in the ovary during the gonadotropl
J. Biol. Chem. 269:11707-11713(1994).
-!- FUNCTION: This enzyme condenses acetyl-Co
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Blattella germanica (German cockroach).
Blattella germanica (German cockroach).
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta;
Neoptera; Orthopteroidea; Dictyoptera; Blattaria;
Blattellidae; Blattellinae; Blattella.
                                                                                                                                                                                                                                                                                      This SWISS-PROT entry is copyright. It is produced through a between the Swiss Institute of Bioinformatics and the EMBI the European Bioinformatics Institute. There are no restrict use by non-profit institutions as long as its content is modified and this statement is not removed. Usage by and fentities requires a license agreement (See http://www.isb-sikor send an email to license@isb-sib.ch).
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                                                                                                                                                Local Sines 246;
                                                                                                                                                                                                                                                                                                                                                                                similarity).

CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acet hydroxy-3-methylglutaryl-CoA + CoA.

PATHWAY: PRODUCTION OF MEVALONATE FROM HMG-CO SYNTHESIS OF STEROLS AND ISOPRENOIDS SUCH AS
                                                                                                                                                                                                                                                                                                                                                                                                                                lated in the ovary during the gonadotrophic cycle iol. Chem. 269:11707-11713(1994). FUNCTION: This enzyme condenses acetyl-CoA with a form HMG-CoA, which is the substrate for HMG-CoA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 TaxID=6973;
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Similarity 53.7%;
16; Conservative 6
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                                                                              TVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDIEGIDTTNACY
                                              GGTAAVFNAVNWIESSSWD-
RĠTAALFNALIWIESSSWDGRYAIVVAADIAIYAKECSPTGGAGALLMLIGANAPIVIDR
                                                                   HWPEDVGIIGIEMIFPSLYVDQAELETYDEVSPGKYTMGLGQDKMGVCTDREDINSLCL
                                                                                                                              ACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCTDREDINSLCM
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cy during the gonadotrophic cyc
                                                                                                                                                    64;
                                                                                                                                                  Score 1272.5; DB 1; Pred. No. 2.2e-85; ; Mismatches 101;
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Blaberoidea;
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CSTRAIN=CV. COlumbia;

X MEDLINE=20083488; PubMed=10617198;

X MEDLINE=20083488; PubMed=10617198;

X Mayer K.F.X., Schueller C., Wambutt R., Murphy G., Volckaert G.,

A Pohl T., Duesterhoeft A., Stiekema W., Entian K.-D., Terryn N.,

A Harris B., Ansorge W., Brandt P., Grivell L.A., Rieger M.,

A Weichselgartner M., de Simone V., Obermaier B., Mache R., Mueller M.,

A Kreis M., Delseny M., Puigdomenech P., Watson M., Schmidtheini T.,

A Kreis M., Delseny M., Puigdomenech P., Watson M., Schmidtheini T.,

A Kreis M., Delseny M., Puigdomenech P., Watson M., Schmidtheini T.,

A Kreis M., Delseny M., Puigdomenech P., Watson M., Schmidtheini T.,

A Kreis M., Delseny M., Perez-Alonso M., Boutry M., Bancroft I.,

A Kreis M., Delseny M., Wedler H., Robben J.,

A Kreis M., Delseny M., Wedler H., Robben J.,

A Kreis M., Defoor E.,

A Weitzenegger T., Bothe G., Ramsperger U., Hilbert H., Braun M.,

A Weitzenegger T., Bothe G., Ramsperger U., Hilbert H., Braun M.,

A Mooijman P., Klein Lankhorst R., Rose M., Hauf J., Koetter P.,

A Mooijman P., Klein Lankhorst R., Rose M., Hauf J., Koetter P.,

A Berneiser S., Hempel S., Feldpausch M., Lamberth S., Van den Daele H.,

A De Keyser A., Buysshaert C., Gielen J., Villarroel R., De Clercq R.,

RA Van Montagu M., Rogers J., Cronin A., Quail M.A., Bray-Allen S.,

RA Pettett A., Rajandream M.A., Lyne M., Lennard N., McLay K., Mayes R.,

RA Pottett A., Rajandream M.A., Lyne M., Brens V., Rechmann S.,

RA Dose S., de Haan M., Maarse A.C., Schaefer M., Mueller-Auer S.,

RA Gabel C., Fuchs M., Fartmann B., Granderath K., Dauner D., Herzl A.,
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P54873; Q8L721; Q9S707;

01-OCT-1996 (Rel. 34, Created)

28-FEB-2003 (Rel. 41, Last sequence update)

15-MAR-2004 (Rel. 43, Last annotation update)

Hydroxymethylglutaryl-CoA synthase (EC 2.3.3.10) (HMG-CoA synthase).

(3-hydroxy-3-methylglutaryl coenzyme A synthase).

HMGS OR MVA1 OR AT4G11820 OR T26M18.30.

Arabidopsis thaliana (Mouse-ear cress).

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta, Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosic eurosids II; Brassicales; Brassicaceae; Arabidopsis.

NCBI_TaxID=3702;
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Connolly E.L., Learned R.M.;
"Post-transcriptional regulation of HMG-CoA synthase expanded and the control of the control of
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STRAIN=cv. Landsberg erecta;
MEDLINE=96144274; PubMed=8566777;
Montamat F., Guilloton M., Karst F., Delrot S.;
"Isolation and characterization of a cDNA encoding Arabidopsis thaliana 3-hydroxy-3-methylglutaryl-coenzyme A synthase.";
Gene 167:197-201(1995).
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r. Columbia;
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Clercq R.,
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A Massenet O., Quigley F., Clabauld G., Muendlein A., Felber R.,
A Massenet O., Quigley F., Clabauld G., Muendlein A., Felber R.,
Schnabl S., Hiller R., Schmidt W., Lecharny A., Aubourg S.,
A Chefdor F., Cooke R., Berger C., Monfort A., Casacuberta E.,
A Gibbons T., Weber N., Vandenbol M., Bargues M., Terol J., Torres A.,
Perrez-Perez A., Purnelle B., Bent E., Johnson S., Tacon D., Jesse T.
A Heijnen L., Schwarz S., Scholler P., Heber S., Francs P., Bielke C.,
A Frishman D., Haase D., Lemcke K., Mewes H.-W., Stocker S.,
A Parnell L., Dedhia N., Gnoj L., Schutz K., Huang E., Spiegel L.,
A Stoneking T., Kalicki J., Graves T., Harmon G., Edwards J.,
Stoneking T., Kalicki J., Graves T., Harmon G., Edwards J.,
A Stoneking T., Kalicki J., Graves T., Harmon G., Edwards J.,
A Kramer J., Fulton B., Miller N., Greco T., Kemp K.,
A Nelson J., Spieth J., Mardis E., Dante M., Pepin K., Hillier L.W.,
A Nelson J., Spieth J., Brandis E., Dante M., Pepin K., Hillier L.W.,
A Antonoiu B., Zidanic M., Strong C., Sun H., Lamar B., Yordan C.,
A Antonoiu B., Zidanic M., Strong C., Sun H., Lamar B., Yordan C.,
A Antonoiu B., Zidanic M., Strong C., Sun H., Lamar B., Yordan C.,
A Antonoiu B., Zidanic M., Strong C., Sun H., Lamar B., Yordan C.,
A Mas P., Zhong J., Preston R., Vil D., Shekher M., Matero A., Shah R.,
A Granat S., Shohdy N., Haeegawa A., Hameed A., Lodhi M., Johnson A.,
A Granat S., Shohdy N., Haeegawa A., Hameed A., Lodhi M., Johnson A.,
A Granat S., Shohdy N., Haeegawa A., Hameed A., Lodhi M., Johnson A.,
A Granat S., Shohdy N., Haeegawa A., Hameed A., Lodhi M., Johnson A.,
A Granat S., Shohdy N., Haeegawa A., Hameed A., Lodhi M., Johnson A.,
A Granat S., Shohdy N., Haeegawa A., Hameed A., Lodhi M., Johnson A.,
B Seguence and analysis of chromosome 4 of the plant Arabidopsis
T. Thailana.",
B Seguence and analysis of chromosome 4 of the plant Arabidopsis
T. Thailana. ",
B Seguence A., B Shohdy N., B Shohdy R., B Shohdy R
                                        EMBL; X83882; CAA58763.1; -.
EMBL; U79160; AAD00297.1; -.
EMBL; U79161; AAD00298.1; -.
EMBL; AL078606; CAB44320.1; -.
EMBL; AL161532; CAB78225.1; -.
EMBL; AY140008; AAM98150.1; -.
EMBL; BT008492; AAP37851.1; -.
PIR; T09341; T09341.
InterPro; IPR000590; HMG_CoA_s
InterPro; IPR008260; HMG_CoA_s
Pfam; PF01154; HMG_CoA_synt; 1
PROSITE; PS01226; HMG_COA_SYNT
Sterol biosynthesis; Transfera
ACT_SITE 117 117
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EMBL; U79160; AAD00297.1; -.

EMBL; U79161; AAD00298.1; -.
  CONFLICT
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- CATALYTIC ACTIVITY: Acetyl-CoA +
hydroxy-3-methylglutaryl-CoA + C
- PATHWAY: Production of mevalonat
synthesis of sterols and isopren
- ALTERNATIVE PRODUCTS:
Event=Alternative splicing; Name
Name=1;
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IsoId=P54873-2; Seq
SIMILARITY: Belongs to
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s to the HMG-C
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and isca--
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POTENTIAL.

Missing (in isoform;
/FTId=VSP_008902.
A -> S (IN REF. 1).
K -> N (IN REF. 1).
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RESULT 12

HMCS YEAST
ID 10-CCT-1996 (Rel. 34, Created)
DT 01-CCT-1996 (Rel. 34, Last sequence update)
DT 01-CCT-1996 (Rel. 34, Last sequence update)
DT 10-CCT-2093 (Rel. 42, Last annocation update)
DT 10-CCT-2093 (Rel. 42, Last sequence update)
DT 10-CCT-2093 (Rel. 42, Last sequence A synthase)
DT 10-CCT-2093 (Rel. 42, Last sequence A synthase)
DT 10-CCT-2093 (Rel. 42, Last sequence as ynthase)
DT 10-CCT-2093 (Rel. 42, Last sequence as ynthase)
DT 10-CCT-2093 (Rel. 42, Last sequence as ynthase)
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runction: This enzyme condenses acetyl-CoA with acetoacetyl-CoA form HMG-CoA, which is the substrate for HMG-CoA reductase (By similarity).

CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acetoacetyl-CoA = (S)-2
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EMBL; X96617; CAA90557.1; -.
EMBL; Z50178; CAA90557.1; -.
PIR; S58202; S58202.
GermOnline; 142661; -.
SGD; S0004595; ERG13.
InterPro; IPR000590; HMG_COA_synt_AS.
InterPro; IPR008260; HMG_COA_synt.
Pfam; PF01154; HMG_COA_synt; 1.
PROSITE; PS01226; HMG_COA_SYNTHASE; 1.
Transferase; Sterol biosynthesis.
ACT_SITE 159 159 POTENTIAL.
SEQUENCE 491 AA; 55013 MW; 44DFF3C0B0
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HMCS SCHPO STANDARD; PRT; 447 AA.
P54874;
01-OCT-1996 (Rel. 34, Created)
01-OCT-1996 (Rel. 34, Last sequence update)
10-OCT-2003 (Rel. 42, Last annotation update)
Hydroxymethylglutaryl-CoA synthase (EC 2.3.3.10)
(3-hydroxy-3-methylglutaryl coenzyme A synthase)
HCS OR SPAC4F8.14C.
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PATHWAY: Production of mevalonate from HMG-CoA prio
synthesis of sterols and isoprenoids.
SIMILARITY: Belongs to the HMG-CoA synthase family.
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EMBL; U32187; AAB17601.1; -.
EMBL; Z98530; CAB11060.1; -.
PIR; S61875; S61875.

GeneDB_SPombe; SPAC4F8.14C; -.
InterPro; IPR000590; HMG_CoA_synt_AS.
InterPro; IPR008260; HMG_CoA_synth.
Pfam; PF01154; HMG_COA_synt; 1.
PROSITE; PS01226; HMG_COA_SYNTHASE; 1.
Transferase; Sterol biosynthesis.

Transferase;
ACT_SITE 1
SEQUENCE 44

118 447 *i*

49239 MW;

POTENTIAL

919BDDBD9207B886

CRC64;

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MEDLINE-21844401; PubMed=11859360;

MEDLINE-2184401; PubMed=11859360;

Mood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,

Mood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,

A Sgouros J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,

A Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,

A Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,

A Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,

A Holroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagels K.,

A James K., Jones M., Leather S., McDonald S., McLean J.,

A Holroyd S., Hornsby T., Howarth S., McDonald S., McLean J.,

A Holroyd S., Hornsby T., Howarth S., McDonald S.,

A Hollower K., O'Neil S., Pearson D., Quail M.A., Rabbinowitsch E.,

A Hollower K., Gowle S., Mungall K., Murphy L., Niblett D., Odell C.,

A Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,

A Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,

A Skelton J., Simmonds M., Squares R., Schaefer M., Whitehead S.,

A Weltjens I., Vanstreels E., Rieger M., Schaefer M., Whitehead S.,

A Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mutler-Auer S.,

A Weltjens I., Vanstreels E., Rieger M., Schaefer M., Mutler-Auer S.,

A Weltjens I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,

Begr P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,

Begry M., Langer I., Beck A., Lehrach H., Reinhardt R., Pohl T.M.,

A Gaffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,

A Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,

A Goffeau A., Rochet M., Gaillardin C., Tallada V.A., Garzon A., Thode G.,

A Dominguez A., Parts J., Jimenz J., Sanchez M., del Rey F., Benito J.,

A Dominguez A., Parts J., Jimenz J., Sanchez M., del Rey F., Benito J.,

A Dominguez A., Parts J., Jimenz J., Sanchez M., del Rey F., Benito J.,

A Dominguez A., Parts J., Jimenz J., Sanchez M., Jenschurg P.,

"The genome sequence of Schizosacharomyces pombe.";

The Grand H., Charles 
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Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;

Schizosaccharomycetales; Schizosaccharomycetaceae;

Schizosaccharomyces.

NCBI_TaxID=4896;

[1]

SEQUENCE FROM N.A.

MEDLINE=96353436; PubMed=8750242;

MEDLINE=96353436; PubMed=8750242;

MEDLINE=96353436; PubMed=8750242;

MEDLINE=96353436; PubMed=8750242;

MEDLINE=96353436; PubMed=8750242;

MEDLINE=96353436; PubMed=11859360;

MEDLINE=21848401; PubMed=11859360;

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ydroxy-3-methylglutaryl coenzyme A synthase of Schizosaccharomyces
oombe.";
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CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acetoacetyl-CoA = CATALYTIC ACTIVITY: Acetyl-CoA + CoA.

hydroxy-3-methylglutaryl-CoA + CoA.

pathway: Production of mevalonate from HMG-CoA prior to the pathway: Production of mevalonate from HMG-CoA prior to the synthesis of sterols and isoprenoids.

SIMILARITY: Belongs to the HMG-CoA synthase family.
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Q22962;

01-OCT-1996 (Rel. 41, Last sequence upd

10-OCT-2003 (Rel. 42, Last annotation upd

10-OCT-394) to the EMBL/GenBan

10-OCT-1994) to the EMBL/GenBan

11-FUNCTION: This enzyme condenses ace

11-OCT-1994) to the EMBL/GenBan

11-OCT-1994) to the EMBL/GenBan
form HMG-con, similarity).

- CATALYTIC ACTIVITY: Acetyl-CoA + H(2) hydroxy-3-methylglutaryl-CoA + CoA. l-PATHWAY: Production of mevalonate from synthesis of sterols and isoprenoids synthesis of sterols and isoprenoids l-SIMILARITY: Belongs to the HMG-CoA sterols.
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Matches 57
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Submitted (MAR-1996) to the EMBL/GenBank/DDBJ databases.
-!- FUNCTION: This enzyme condenses acetyl-CoA with acetoacetyl-CoA to form HMG-CoA, which is the substrate for HMG-CoA reductase (By similarity).
-!- CATALYTIC ACTIVITY: Acetyl-CoA + H(2)O + acetoacetyl-CoA = (S)-3-hydroxy-3-methylglutaryl-CoA + CoA.
-!- PATHWAY: Production of mevalonate from HMG-CoA prior to the synthesis of sterols and isoprenoids.
-!- SIMILARITY: Belongs to the HMG-CoA synthase family.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See http://www.isb-sib.ch/announce/or send an email to license@isb-sib.ch).
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protein cDNA.

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isolated nucleic acid molecule encoding hydroxymethylglutaryl-CoAthase, useful as model for the development of human therapeutic gets and for identifying therapeutic proteins.
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SDB; ABG32726.
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standard; cDNA; 3722

ADE76933;

29-JAN-2004

Human cDNA differentially expressed in (first entry) a liver disorder #70

human; ss; gene; liver disorder; hyperlipidaemia; hypertension; type II diabetes; tumour; liver; inflammatory disorder; immune response disorder; high-throughput screening; differential gene expression; gene therapy.

Homo sapiens.

US2003108871-A1

12-JUN-2003

30-JUL-2001; 2001US-00919039.

28-JUL-2000; 2000US-0222113P

KASER ™ 72.

Kaser MR ;

WPI; 2004-031227/03

Composition comprising several cDNAs that treated human C3A liver cell cultures, use disorders. nat are differentially expressed useful for treating liver

Claim SEQ ID NO 98; 41pp; English

The invention relates to a composition comprising several cDNAs that are differentially expressed in a liver disorder. The composition is useful for treating liver disorder such as hyperlipidaemia, hypertension, type II diabetes, tumours of the liver and disorders of the inflammatory and immune response. The composition is useful for a high-throughput method of screening several molecules or compounds to identify a ligand which specifically binds a cDNA. A protein encoded by the cDNA is useful for a high-throughput method for using a protein to screen several molecules or compounds to identify at least one ligand which specifically binds the protein which involves combining the protein encoded by the cDNA with several of molecules or compounds under conditions to allow specific binding, and detecting specific binding between the protein and a og a

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CE, Ooi CE, Ort Rothenberg ME, KA, Taupier RJ,

Giot L, Gorman L, Guo X, Gusev VY, Ji W Leach MD, Lepley DM, Li L, Liu X; Ooi CE, Ort T, Padigaru M, Patturajan M; thenberg ME, Shenoy SG, Shimkets RA; Taupier RJ, Twomlow N, Vernet CAM, Voss E

Voss EZ;

gee ML, Alsobrook JP, Aringess CE, Casman SJ, Carabtree J, Dipippo VA, Ingolli EA, Gerlach VL, kuda R, Khramtsov NV, Inlyankar UM, Miller CE,

Anderson DW,
Catterton E,
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Charcacteristic anticological in the exemplification of the present invention with the human NOV35b protein in the exemplification of the present invention.
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                                                                                           Agee ML, Alsobrook JP, Anderson DW, IBurgess CE, Casman SJ, Catterton E, Crabtree J, Dipippo VA, Edinger SR, IGangolli EA, Gerlach VL, Giot L, Gorr Kekuda R, Khramtsov NV, Leach MD, Lemalyankar UM, Miller CE, Ooi CE, Ort Pena CEA, Rieger DK, Rothenberg ME, Spaderna SK, Spytek KA, Taupier RJ, Zerhusen BD, Zhong M;
     Novel human diagnosis,
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P-PSDB; ABR54259.
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2001US-0323519P.
2001US-0323631P.
2001US-0323631P.
2001US-0323631P.
2001US-0324969P.
2001US-0324990P.
2001US-0341144P.
2002US-0361663P.
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PT protein or nucleic acid e.g. cardiac and neurological disorders.

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Claim 20; Page 241; 460pp; English.

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CC The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in CC ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, CC Appotensive, dermatological, anorectic, immunosuppressive, cycostatic, CC antidiabetic, antiinfertility, haemostatic, neuroprotective, notropic, CC antiparkinsonian and antilipaemic activities, and can be used in gene CC therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome CC associated with the human disease. NOVX nucleic acids, proteins and CC antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, disbetes, metabolic disease, tuberous sclerosis, scleroderma, obesity, transplantation, CC disorders, nooplasm, lymphoma, uterus cancer, fertility, haemophilia, hyperplasia, prostate cancer, disbetes, metabolic disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, hearting cancer associated cachexia, cancer, ACC62346 to ACC6245 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention.

CC ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention.
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Sequence 1650 BP; 477 A; 319 C; 397 G; 457 T; 0 U; 0 Other;

Query Match
68.4%; Score 1370; DB 7; Length 1650;
Best Local Similarity 92.0%; Pred. No. 4.1e-259;
Matches 1509; Conservative 0; Mismatches 5; Indels 126; Gaps 1;
Qy 134 GCTCTTTCACCATGCCTGGATCACTTCCTTTGAATGCAGAAGCTTGCTGGCCAAAAGATG 193

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CAGATATAGAAGGAATCGACACAACTAATGCATGCTATGGAGGCACAGCTGCTGTCT 553	ATACA	494	γQ
TCGACAAATCAAAGTCTGTGAAGACTAATTTGATGCAGCTGTTTGAAGAGTCTGGGA 370	Ω= C2=	311	DЬ
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TCTGCACAGATAGAGAAGATATTAACTCTCTTTGCATGACTGTGGTTCAGAATCTTA 373	GCTTC	314	γQ
AAAAATATGATGGTGTAGATGCTGGGAAGTATACCATTGGCTTGGGCCAGGCCAAGATGG 190		131	DЬ
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TTGGGATTGTTGCCCTTGAGATCTATTTTCCTTCTCAATATGTTGATCAAGCAGAGTTGG 130	TTGGG	71	Дb
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ACC62326 standard;
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Homo sapiens

WO2003023001-A2

20-MAR-2003

07-SEP-2001; 10-SEP-2001; 17-SEP-2001; 17-SEP-2001; 17-SEP-2001; 17-SEP-2001; 19-SEP-2001; 20-SEP-2001; 20-SEP-2001; 25-SEP-2001; 25-SEP-2001; 26-SEP-2001; 26-FEB-2002; 05-MAY-2002; 03-MAY-2002; 17-JUL-2002; 13-AUG-2002; 13-AUG-2002; 09-SEP-2002; 2001US-0318120P.
2001US-0318120P.
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2002US-0393332P.
2002US-0396412P.
2002US-0403517P.
2002US-0403517P. 2002WO-US028538

(CURA-) CURAGEN CORP

Agee ML, Alsobrook JP, Anderson DW, Berghs C, Boldog FL;
Burgess CE, Casman SJ, Catterton E, Chant JS, Chaudhuri A;
Crabtree J, Dipippo VA, Edinger SR, Eisen AJ, Ellerman K;
Gangolli EA, Gerlach VL, Giot L, Gorman L, Guo X, Gusev VY, Ji W
Kekuda R, Khramtsov NV, Leach MD, Lepley DM, Li L, Liu X;
Malyankar UM, Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M;
Pena CEA, Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA;
Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss E;
Zerhusen BD, Zhong M; Voss EZ; Ξ

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Agee ML, Alsobrook is Burgess CE, Casman & Crabtree J, Dipippo Gangolli EA, Gerlach Kekuda R, Khramtsov Malyankar UM, Miller Pena CEA, Rieger DK, Spaderna SK, Spytek Zerhusen BD, Zhong Manger B
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Chaudhuri A;
Ellerman K;
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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antialabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NoVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital edisease, tuberous sclerosis, scleroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host chiesese, anorexia, cancer-associated cachexia, cancer, halps, bronchial asthma, Crohn's disease, immune disorders, cancer, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. Acc62346 to ACC62465 represent PCR primers and probes for human NOVX seguences, which are used in examples from the present invention.

Charleting the human novasb protein in the exemplification of the present invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Novel human proteins and nucleic acid encoding the proteins, useful diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.
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P-PSDB; ABR54261.
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2001US-0318130P.
2001US-0318430P.
2001US-0322636P.
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Catterton E,
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Leach MD, I n E, Gorman L Lepley | Ort T, Berghs C, Chant JS, Eisen AJ, DM, Padigaru JS, Guo Boldog FL; Chaudhuri A; Ellerman K; no X, Gusev V Li L, Liu X; garu M, Pattu Gusev V Liu X; Patturajan M; A; ٧٧, Ji

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Pena CER. Rieger DK. Rethenberg ME. Schenov MG. Schinchets RA. Spederma SK. Spytek KA. Taupker RE. Twomlow N. Vernet CAM. Vose EZ. Zechusen BD. Zhong N. WPI: 2003-313241/30. P-PESDR, Amstages. Noval human proteins and mucleic acid encoding the proteins, where X is a chimacosts, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders. Claim 20; Page 246, 460pp. English. The prosent invention describes incole the human NOWX proteins, where X is 1 to 42. Co56223 to Accessable encode the human NOWX proteins of the cardiac and neurological disorders. ABSELIC: antilideric antilingericity, hereosetatic, ministaberosclerotic, cardiant, hypotensive, dermacological, anorectic, immunosuppressive, optensation, antidaberic, antilingericity, hereosetatic, ministaberoscleroty, anti-sity, antidaberic, dermacological, anorectic, immunosuppressive, cytestarid, hypotensive, dermacological, anorectic, immunosuppressive, cytestarid, antibodies can be used in the treatment and disgnosts of cardiant, hypotensive, thermacological, anorectic, and proventing a pathony associated with a KONY protein in humans and for treating a pathony associated with a KONY protein in humans and for treating a pathony associated with a KONY protein in the access and protein contexts, cancer, anorectic, antipathy and the protein and antibodies can be used in the treatment and disgnosts of cardinary hypoteosety, protein and anorexia, cancer associated cachests, cancer, hypoteosety, protein and protein protein and anorexia, cancer, hypoteosety, protein and anorexia, cancer associated cachests, cancer, hypoteosety, protein and anorexia, cancer-associated cachests, cancer, hypoteosety, protein and the acceptance and anothers, and the respective protein and hypoteo
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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, valve disease, tuberous sclerosis, scleroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, infectious disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, Parkinson's disease, immune disorders, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X.

ACC62346 to ACC62465 represent PCR primers and probbes for human NOVX sequences, which are used in examples from the present invention.

ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        The present in to 42. ACC6 ABR54167 to A hypotensive, antidiabetic, antiparkinson therapy. NOVX associated with associated with associated with antibodies cantibodies cantibodies; antibodies cantibodies; necongenital addisorders, nethypercoagulations infectious disparents in Alzheimer's dhaematopoietial ACC62346 to A sequences, whalsheimer's dhaematopoietial ACC62346 to A sequences, whalsheimer's disparents whalsheimer's dhaematopoietial ACC62346 to A sequences, whalsheimer's disparents whalsheimer's dhaematopoietial ACC62346 to A sequences, whalsheimer's disparents which was a sequence of the control of the 
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a human NOV35b protein i
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kuda R, Khramtsov NV, I
lyankar UM, Miller CE,
na CEA, Rieger DK, Roth
baderna SK, Spytek KA, T
rhusen BD, Zhong M;
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TCATCGACAAATCAAAGTCTGTGAAGACTAATTTGATGCAGCTGTTTGAAGAGTCTGGGA
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CE, Ooi CE, Ort T, Padigaru M, Patturajan M;
Rothenberg ME, Shenoy SG, Shimkets RA;
A, Taupier RJ, Twomlow N, Vernet CAM, Voss E
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                                                      07-SEP-2001;

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25-SEP-2001;

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Novel human proteins and nuclair acid encoding the proteins, useful for diagnosis, treatment and prevention of disporders involving the human protein or muclair acid e.g. cardiac and neurological disorders. Claim 20; Page 240-241, 46pp; English. The present invention describes isolated human NOVX proteins years in ABS3167 to ABS4276. NOVX sequences have anniable proteins given in ABS3167 to ABS4276. NOVX sequences have anniable proteins given in ABS3167 to ABS4276. NOVX sequences have anniable received the proteins given in ABS3167 to ABS4276. NOVX sequences have anniable received the proteins given in ABS3167 to ABS4276. NOVX sequences have anniable received the proteins given in ABS3167 to ABS4276. NOVX sequences have anniable received the proteins given in ABS3167 to ABS4276. NOVX sequences have anniable received the proteins given in the treatment and for treating any developed, which a NOVX protein in humans and for treating a price of the proteins given the proteins given the proteins given in the treatment and diagnosis of cardiamy selections are all propriates, proteins and for treating a price of the proteins given in the proteins. Crohn staces, diabetes, metabolic disorders, neeplasm, Jymphoma, uterus cancer, fertility, haemophilia, hypercoaghi, cancer associated carbania, cancer subscited carbania granter of disease, annual disorders, metabolic proteins given in concerns an interest of disease, annual disorders, metabolic proteins given in the present invention of the present invention of the human NOV35b protein in the exemplia feating given in comparison with the human NOV35b protein in the exemplia feating given in comparison with the human NOV35b protein in the exemplia feating given in comparison with the human NOV35b protein in the exemplia feating given in comparison with the human NOV35b protein in the exemplia feating given in comparison with the human NOV35b protein given in compa	Agee ML, Alsobrook JP, Anderson DW, Berghs C, Boldog FL; Burgess CE, Casman SJ, Catterton E, Chant JS, Chaudhuri A; Crabtree J, Dipippo VA, Edinger SR, Eisen AJ, Ellerman K; Gangolli EA, Gerlach VL, Giot L, Gorman L, Guo X, Gusev VY, Ji W; Kekuda R, Khramtsov NV, Leach MD, Lepley DM, Li L, Liu X; Malyankar UM, Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M; Pena CEA, Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA; Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; Zerhusen BD, Zhong M;
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Agee ML, Alsobrook JP, Anderson DW, Berghs C, Boldog FL; PI Burgees CE, Casman SJ, Catetron E, Chant JS, Chaudhuri A; Brigges CE, Casman SJ, Catetron E, Chant JS, Chaudhuri A; PI Carbtree J, Djippo VA, Edinger SR, Eisen AJ, Ellerman K; PI Carbtree J, Djippo VA, Edinger SR, Eisen AJ, Ellerman K; PI Catetron E, Cornan L, Guo X, Guev VV, Ji W; Catet MU, Ciot L, Gornan L, Guo X, Guev VV, Ji W; Kokuda R, Khramteov NV, Leach MD, Lepley DM, Li L, Liu X; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; Szerhusen BD, Zhong M; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EZ; PI Spadern

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Wanorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility;

Whaemostatic; antiinflammatory; antiasthmatic; anti-HIV; immunomodulator;

What cancer in control is antiasthmatic; anti-HIV; immunomodulator;

What cancer; antiinflammatory; antiasthmatic; anti-HIV; immunomodulator;

What cancer; metabolic disorder; neoplasm; hyperplasia; diabetes;

What is a control is antiplammatory; and is antimatic; antiplammatory;

What cancer is antiinflammatory; antiasthmatic; antiinflammatory;

What is a control is antiinflammatory; antiasthmatic; antiinflammatory;

What is a cancer; antiinflammatory; antiasthmatic; antiinflammatory;

What is antiilipaemic;

What is anti-HIV; immunomodulator;

gee ML, Alsobrook JP, An Burgess CE, Casman SJ, Ca Crabtree J, Dipippo VA, I Gangolli EA, Gerlach VL, Kekuda R, Khramtsov NV, I Malyankar UM, Miller CE, Pena CEA, Rieger DK, Roth
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10-SEP-2001;

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25-SEP-2001;

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2001US-0322636P

2001US-0322781P

2001US-0322816P

2001US-0323819P

2001US-0323631P

2001US-0323631P

2001US-0323631P

2001US-0324969P

2001US-0324969P

2001US-0324969P

2001US-0341144P

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TEXE EXELECTION OF SOLUTION OF The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in C ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiatherosclerotic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, replasm, lymphoma, uterus cancer, diabetes, metabolic disease, allow, idiopathic thrombocytopenic purpura, graft versus host confectious disease, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, hamman source associated cachexia, cancer, anorexia, cancer-associated cachexia, cancer, caches, which are used in examples from the present invention.

CC ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention. Spaderna Zerhusen WPI; 2003~313241/30.
p-PSDB; ABR54268.

Novel human proteins and nucleic acid encoding the proteins, useful diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders. SK, 1564 Spytek Zhong N BP; 245-246; 460pp; English. 3 Ķ¥, Taupier <u>ე</u> 362 G; 434 T; 0 U; 0 Other; Twomlow N, Vernet CAM, Voss for EZ;

Š Š 밁 Ş 밁 멍 Ş В Ş 멍 80 Š 망 Query Match Best Local S Matches 1438 al Similarity 144 181 121 264 204 241 564 361 61 Н TGCCCTTGAGATCTATTTTCCTTCTCAATATGTTGATCAAGCAGAGTTGGAAAAATATGA CATGCCTGGATCACTTCCTTTGAATGCAGAAGCTTGCTGGCCAAAAGATGTGGGAATTGT ĊĄŦĠĊĊŦĠĠĄŦĊĄĊŦŦĊĊŦŦŦĠĄĄŦĠĊĄĠĄĄĠĊŦŦĠĊŦĠĠĊĊĄĄĄĄĠAŦĠŦĠĠĠĄĄŦŦĠŦ TGGTGTAGATGCTGGAAAGTATACCATTGGCTTGGGCCAGGCCAAGATGGGCTTCTGCAC TAACTGGATTGAGTCCAGCTCTTGGGAT-65.0%;
.larity 91.9%;
Conservative 0; Score Pred. ore 1302; DB 7; ed. No. 8.6e-246; Mismatches 0; Indels 126; Gaps 203 503 180 263 09 563 360 300 443 240 383 323 120 420 591 بر ،،

1577 1560	1518 ACCAAGACTCCCTGCCACAGCAGCAGAACCTGAAGCAGCTGTCATTAGTAATGGGGAACA
1517 1500	58 AGTAGGACTTGTGCATTCAAACATAGCAACTGAGCATATTCCAAGCCCT
1457 1440	1398 AAAGCACAGAAGAACTTACGCTCGGCGTCCCACTCCAAATGATGACACTTTGGATGAAGG
1397 1380	1338 TCCCCAGGGTTCAATAGATTCACTCTTTGAAGGAACGTGGTACTTAGTTAG
1337 1320	1278 AGATGTCTTCGCTGAAAACATGAAGCTCAGAGAGAGACACCCATCATTTGGTCAACTATAT
1277 1260	1218 AATAACAGCAAGTTTATGTGATCTTAAATCAAGGCTTGATTCAAGAACTGGTGTGGCACC
1217	1158 TGCCACTCTGTACTCTCTTAAAGTCACACAAGATGCTACACCGGGGTCTGCTCTTGATAA
1157 1140	1098 CTCACCTCAGCAATTAGCAGGGAAGAGAATTGGAGTGTTTTCTTATGGTTCTGGTTTTGGC
1097	1038 AAATGGAAATATGTACACATCTTCAGTATATGGTTCCCTTGCATCTGTTCTAGCACAGTA
1037	978 GAAGGCTAGCTCTGAACTCTTCAGTCAGAAAACAAAGGCATCTTTACTTGTATCAAATCA
977 960	918 CTTTGGGGATGTTAAATTAGAAGACACCTACTTTGATAGAGATGTGGAGAAGGCATTTAT
917 900	858 GAATGACTTCCTTAATGACCAGAATAGAGATAAAAATAGTATCTATAGTGGCCTGGAAGC
857 840	798 CATGATCTTTCACTCACCATATTGTAAACTGGTTCAGAAATCTCTAGCTCGGATGTTGCT
797 780	738 CCATGCCCAGTGGCAGAAAGAGAGGGAAATGATAAAGATTTTACCTTGAATGATTTTGGCTT
737 720	678 CTCCATACAGTGCTACCTCAGTGCATTAGACCGCTGCTACTCTGTCTACTGCAAAAAGAT
677	618 TGCCTATGATTTTTACAAGCCTGATATGCTATCTGAATATCCTATAGTAGATGGAAAACT
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Sequence 9, Appli	LU -	Sequence 10, Appl	ce 67,	•	, App	equence 24, A	equence 22, App	4, App	equence 22, App	equence 2	0	equence 8	equence 8, App		6, App	Sequence 12, Appl	

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; Patent No. 6620608
; GENERAL INFORMATION:
APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND
TITLE OF INVENTION: THEREOF
FILE REFERENCE: CL001195DIV
CURRENT APPLICATION NUMBER: US/10/193,295
; CURRENT FILING DATE: 2002-07-12
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pRIOR FILING DATE: 2001-03-29
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FASTSEQ for Windows Version 4.0
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                                                AGCGAAGGGGAGGCGCCGCGGACTGTCCTTTCGTGGCTCACTCCCTTTCCTCTGCTGCCG
    CTCGGTCACGCTTGCTCTTCACCATGCCTGGATCACTTCCTTTGAATGCAGAAGCTTGC
                                                                                 0;
                                                                                Score 2002;
Pred. No. 0;
0; Mismatches
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                                                                                                Length
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US-08-305-505-1

; Sequence 1, Application US/08305505

; Patent No. 5668001

; Patent No. 5668001

; GENERAL INFORMATION:
  APPLICANT: Miziorko, Henry M.
  TITLE OF INVENTION: 3-HYDROXY-3-METHYLGLUTARYL-COA

; TITLE OF INVENTION: SYNTHASE PREPARATION WITH IMPR

; TITLE OF INVENTION: STABILITY

NUMBER OF SEQUENCES: 6

; CORRESPONDENCE ADDRESS:
  ADDRESSEE: Quarles & Brady
  STREET: 411 East Wisconsin Avenue
  CITY: Milwaukee
  STATE: Wisconsin
  COUNTRY: U.S.A.
  ZIP: 53202

COMPUTER READABLE FORM:
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COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Vel
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/305,505
FILING DATE:
CLASSIFICATION DATA:
APPLICATION NUMBER: US/08/072,040
FILING DATE: 02 JUNE 1993
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 65-053-908:
TELEPHONE: (414) 277-5709
TELEPHONE: (414) 277-5591
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1824 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: CDNA
US-08-305-505-1
                                                                                                                                                                                                                                                                                                                                                                      Query Match 39.7%;
Best Local Similarity 72.3%;
Matches 1139; Conservative
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Pred. No. 4e-185;
0; Mismatches 304;
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RESULT 4
US-09-819-993-3
; Sequence 3, Application US/09
; Patent No. 6436692
; GENERAL INFORMATION:
; APPLICANT: GONG, Fangcheng e

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AGGAGTTGAAGTTGTCCACCCAGGCATTGTTCATGAGCACATCCCAAGCCCTGCTAAGAA	AGGAGT	1441	đđ
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ACCAGATGTCTTCGCTGAAAACATGAAGCTCAGAGAGGACACCCATCATTTGGTCAACTA	ACCAGA]	1275	γQ
	CAAAATA	1201	Db
ACAGCAAGTTTATGTGATCTTAAATCAAGGCTTGATTCAAGAACT	TAAAATA	1215	8
ACGCTGTATTCCATCAGAGTTACACAGGATGCCACTCCTGGTTCTV	TGCTGCT	1141	Db
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CAGTGGCAAAAAGAGGGGACAGACAGAGGTTTCACCTTGAATGATT	CCATGCC	721	Db
CCATGCCCAGTGGCAGAAAGAGGGAAATGATAAAGATTTTACCTTGAATGATTTTGGCTT	CCATGCC	738	Qy
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CAGTGCTACCTCAGTGCATTAGACCGCTGCTACTCTGTCTACTGCA	CTCCATA	678	γQ
${ t TGCTTATGACTTCTATAAACCAGATATGGTTTCTGAATATCCTGTAGTTGATGGCAAACT}$	TGCTTAT	601	Db

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RESULT 5
US-10-193-295-3
; Sequence 3, Application US/10193295
; Patent No. 6620608
; GENERAL INFORMATION:
; APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYR
TITLE OF INVENTION: ACID MOLECULES ENCO
TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001195DIV
; CURRENT APPLICATION NUMBER: US/10/193, 29
; PRIOR APPLICATION NUMBER: 08/819, 993
; PRIOR FILING DATE: 2001-03-29
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LENGTH: 28001
TYPE: DNA
ORGANISM: Human
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(28001)
OTHER INFORMATION: n = 1
US-09-819-993-3
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     TITLE OF INVENTION: ISOLATED HUMAN ENZ
TITLE OF INVENTION: ACID MOLECULES EN
TITLE OF INVENTION: THEREOF
FILE REFERENCE: CLOO1195
CURRENT APPLICATION NUMBER: US/09/819,
CURRENT FILING DATE: 2001-03-29
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSEQ for Windows Version
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Best Local S
Matches 491
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LENGTH: 28001
TYPE: DNA
ORGANISM: Human
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(28001)
OTHER INFORMATION: n = A
US-10-193-295-3
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Similarity 95.2%;
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                                                        GTTGTGGTCTGGCCAATGCCAAATGTACTCGAATGATGTTAAGGGCTCTGTAAAACTTCA
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RESULT 6
US-09-370-838-245
; Sequence 245, Application US/09370838
; Patent No. 6444425
; GENERAL INFORMATION:
APPLICANT: Reed, Steven G.
APPLICANT: Lodes, Michael J.
APPLICANT: Mohamath, Roadoh
APPLICANT: Secrist, Heather
TITLE OF INVENTION: COMPOUNDS FOR THERAPY AND DIAGNOSIS; TITLE OF INVENTION: LUNG CANCER AND METHODS FOR THEIR US
FILE REFERENCE: 210121.475C1
CURRENT APPLICATION NUMBER: US/09/370,838
CURRENT FILING DATE: 1999-08-09
EARLIER APPLICATION NUMBER: US 09/285,323
EARLIER FILING DATE: 1999-04-02
NUMBER OF SEQ ID NOS: 289
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 245
LENGTH: 615
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US-09-370-838-128

Sequence 128, Application US/09370838;
Patent No. 6444425;
GENERAL INFORMATION:
APPLICANT: Reed, Steven G.
APPLICANT: Lodes, Michael J.
APPLICANT: Mohamath, Roadoh
APPLICANT: Secrist, Heather
TITLE OF INVENTION: COMPOUNDS FOR THERAPY
TITLE OF INVENTION: LUNG CANCER AND METHOR
FILE REFERENCE: 210121.475C1;
CURRENT APPLICATION NUMBER: US/09/370,838;
CURRENT FILING DATE: 1999-08-09;
EARLIER APPLICATION NUMBER: US 09/285,323;
BARLIER APPLICATION NUMBER: US 09/285,323;
WUMBER OF SEQ ID NOS: 289
SOFTWARE: FASTSEQ for Windows Version 3.C
SEQ ID NO 128
LENGTH: 500
TYPE: DNA
ORGANISM: Homo sapien
US-09-370-838-128
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ORGANISM: Homo sapie
FEATURE:
NAME/KEY: misc featur
LOCATION: (105)
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09-370-838-245
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APPLICANT: Xu, Jiangchun
APPLICANT: Lodes, Michael J.
APPLICANT: Lodes, Michael J.
APPLICANT: Lodes, Michael J.
APPLICANT: Secrist, Heather
APPLICANT: Secrist, Heather
APPLICANT: Secrist, Heather
APPLICANT: Benson, Darin R.
APPLICANT: Meagher, Madeline Joy
APPLICANT: Wang, Tongtong
TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY AND
TITLE OF INVENTION: DIAGNOSIS OF COLON CANCER AND METH
FILLE REFERENCE: 210121.471C2
CURRENT APPLICATION NUMBER: US/09/401,064
CURRENT FILING DATE: 1999-09-22
NUMBER OF SEQ ID NOS: 371
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 187
LENGTH: 506
TYPE: DNA
ORGANISM: Homo sapien
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US-09-401-0
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Matches 401
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Pred. No. 6.2e-89;
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RESULT 10
US-09-833-381-1648
; Sequence 1648, Application; Patent No. 6672186
; GENERAL INFORMATION:
; APPLICANT: Robison, 1
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NAME/KEY: misc_feature LOCATION: (1)...(448)
OTHER INFORMATION: n = 1-09-833-381-1645
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                                                     NT: Philippsen, Peter
NT: Pohlmann, Rainer
NT: Steiner, Sabine
NT: Mohr, Christine
NT: Wendland, Jurgen
NT: Knechtle, Philipp
NT: Rebischung, Corinne
F INVENTION: GENOMIC DNA S
F INVENTION: AND USES THER
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US/0899841

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LE OF INVENTION: No. 6672186el Nucle REFERENCE: 5800-119
RENT APPLICATION NUMBER: US/09/833, RENT FILING DATE: 2001-04-11
OR APPLICATION NUMBER: 09/516,448
OR FILING DATE: 2000-02-29
BER OF SEQ ID NOS: 2050
TWARE: FastSEQ for Windows Version
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E/KEY: misc_feature
ATION: (1)...(472)
ER INFORMATION: n =
33-381-1648
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RESULT 12
US-09-833-381-744
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STATE: No. 6239264th Carolina
COUNTRY: USA
ZIP: 27709

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENTIN Release #1.0, Ver
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/998,416
FILING DATE: 24-DEC-1997
CLASSIFICATION E 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: CH 0016/97
FILING DATE: 31-DEC-1996
ATTORNEY/AGENT INFORMATION:
NAME: Meigs, J. Timothy
REGISTRATION NUMBER: PF/5-30306/
TELECOMMUNICATION INFORMATION:
TELEPHONE: 919-541-8587
TELEFAX: 919-541-8689
INFORMATION FOR SEQ ID NO: 730:
SEQUENCE CHARACTERISTICS:
LENGTH: 635 base pairs
TYPE: nucleic acid
STRANDEDNESS: Single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORIGINAL SOURCE:
ORGANISM: PAG1476UP
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Best Local Similarity 62.6%;
Matches 231; Conservative
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CURRENT APPLICATION NUMBER: US/09/833,381
CURRENT FILING DATE: 2001-04-11
PRIOR APPLICATION NUMBER: 09/516,448
PRIOR FILING DATE: 2000-02-29
NUMBER OF SEQ ID NOS: 2050
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 744
LENGTH: 307
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NAME/KEY: misc_feature
LOCATION: (1)...(307)

OTHER INFORMATION: n = i
09-833-381-744
                                                                                                                                                                                                                                                                                                                                                                                                                                            TYPE: DNA
ORGANISM: Homo
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Gimilarity 56.9%;
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RESULT 13
US-09-306-595C-1
; Sequence 1, Application US/09306595C
; Patent No. 6284506
; GENERAL INFORMATION:
; APPLICANT: HOSHINO, Tatsuo
APPLICANT: OJIMA, Kazuyuki
APPLICANT: OJIMA, Kazuyuki
APPLICANT: SETOGUCHI, Yutaka
TITLE OF INVENTION: ISOPRENOID PRODUCTION
FILE REFERENCE: ISOPRENOID PRODUCTION
CURRENT APPLICATION NUMBER: US/09/306,595C
CURRENT FILING DATE: 1999-05-06
PRIOR APPLICATION NUMBER: 98108210
PRIOR FILING DATE: 1998-05-06
INUMBER OF SEQ ID NOS: 43
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 1
LENGTH: 4775
TYPE: DNA
ORGANISM: Phaffia rhodozyma
FEATURE:
NAME/KEY: 5'UTR
LOCATION: (1239)..(1240)
OTHER INFORMATION: EXPERIMENTAL
NAME/KEY: exon
LOCATION: (1305)..(1361)

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RESULT 14
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                                                                                    APPLICANT: HOSHINO,
APPLICANT: OJIMA, F
APPLICANT: SETOGUCE
FILE REFERENCE: ISOF
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PPLICANT: OJIMA, Kazuyuki
PPLICANT: SETOGUCHI, Yutaka
PPLICANT: SETOGUCHI, Yutaka
PPLICANT: SETOGUCHI, Yutaka
PRESENCE: ISOPRENOID PRODUCTION
PRESENT APPLICATION NUMBER: US/09/925,388
PRESENT FILING DATE: 2001-08-09
PLIOR APPLICATION NUMBER: 09/306,595
PLIOR FILING DATE: 1999-05-06
PMBER OF SEQ ID NOS: 43
PSTWARE: Patentin Ver. 2.1
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Similarity 63.0%;
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Score 103.6; DB 4;
Pred. No. 1.1e-15;
0; Mismatches 94;
                                                Length
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RESULT 15
US-09-313-294A-4313; Sequence 4313, Application US/09313294A; Patent No. 6476212; GENERAL INFORMATION:
APPLICANT: Lalgudi, Raghunath V.; APPLICANT: Ito, Laura Y.; APPLICANT: Sherman, Bradley K.; TITLE OF INVENTION: POLYNUCLEOTIDES AND POLY FILE REFERENCE: PL-0017 US; CURRENT APPLICATION NUMBER: US/09/313,294A; CURRENT FILING DATE: 1999-05-14; NUMBER OF SEQ ID NOS: 7600; SOFTWARE: PERL Program; SEQ ID NO 4313; LANGTH: 305; TYPE: DNA ORGANISM: Zea mays FEATURE: NAME/KEY: misc feature; NAME/KEY: misc feature; OTHER INFORMATION: Incyte ID No. 6476212 700; US-09-313-294A-4313
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1 2002 100.0 2002 14 US-10-193-295-1 2 2002 100.0 2002 16 US-10-622-516-1 3 1693 84.6 3722 10 US-09-919-039-98 4 1370 68.4 1650 13 US-10-236-417-183 6 1370 68.4 1650 13 US-10-236-417-189 7 1370 68.4 1650 13 US-10-236-417-193 9 1370 68.4 1650 13 US-10-236-417-193 9 1370 68.4 1650 13 US-10-236-417-201 11 1370 68.4 1650 13 US-10-236-417-201 11 1370 68.4 1650 13 US-10-236-417-205 12 1370 68.4 1650 13 US-10-307-817-371 13 1370 68.4 1650 13 US-10-307-817-371 14 1370 68.4 1650 13 US-10-307-817-371	SUMMARIES Query e Match Length DB ID Descr	Pred. No. is the number of results predicted by chance to he score greater than or equal to the score of the result beir and is derived by analysis of the total score distribution.	Database: Published Applications NA:* 1: /cgn2_6/ptodata/2/pubpna/US07_PUBCOMB.seq:* 2: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq:* 3: /cgn2_6/ptodata/2/pubpna/US06_NEW_PUB.seq:* 4: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq:* 5: /cgn2_6/ptodata/2/pubpna/US06_PUBCOMB.seq:* 7: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq:* 9: /cgn2_6/ptodata/2/pubpna/US08_NEW_PUB.seq:* 10: /cgn2_6/ptodata/2/pubpna/US08_PUBCOMB.seq:* 11: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq:* 12: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq:* 13: /cgn2_6/ptodata/2/pubpna/US09B_PUBCOMB.seq:* 14: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq:* 15: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq:* 16: /cgn2_6/ptodata/2/pubpna/US09_NEW_PUB.seq:* 16: /cgn2_6/ptodata/2/pubpna/US10A_PUBCOMB.seq:* 17: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:* 18: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:* 19: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:* 19: /cgn2_6/ptodata/2/pubpna/US10B_PUBCOMB.seq:*	Post-processing: Minimum Match 0% Maximum Match 100% Listing first 45 summaries	Minimum DB seq length: 0 Maximum DB seq length: 20000000000	Total number of hits satisfying chosen parameters: 6034852	Searched: 3017426 segs, 2290544650 residues	Scoring table: IDENTITY_NUC Gapop 10.0 , Gapext 1.0	Title: US-10-622-516-1 Perfect score: 2002 Sequence: 1 cgcctcccagcgactctcggaaaaaaaaaaaaaaa	Run on: June 24, 2004, 09:40:27 ; Search time 840 Seconds (without alignments) 10918.263 Million cell	OM nucleic - nucleic search, using sw model	Copyright (c) 1993 - 2004 Compugen Ltd.
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Sequence 164652,	164	16465	46	Ş	equence 1	Sequence 10, Appl	Sequence 3, Appli	e 3,	Sequence 7, Appli	7,	Sequence 37, Appl	5, App	e 375, A	Sequence 365, App	Sequence 211, App	e 195,	Sequence 367, App	e 357,	e 209,	e 199, A	e 361, A	e 187,	Sequence 363, App	1,	е Э	Sequence 369, App	ce 207,	e 203,	Sequence 1926, Ap	Sequence 55, Appl

ALIGNMENTS

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RESULT 1

US-10-193-295-1

US-10-193-295-1

; Sequence 1, Application US/10193295
; Publication No. US20020173018A1
; GENERAL INFORMATION:
; APPLICANT: GONG, Fangcheng et al.
; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001195DIV
; CURRENT APPLICATION NUMBER: US/10/193, 295
; CURRENT APPLICATION NUMBER: US/10/193, 295
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: 08/819, 993
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 2002
; TYPE: DNA
; ORGANISM: Human
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RESULT 2 US-10-622-516-1 Sequence 1, Application US/10622516 Sequence 1, Application US/10622516 Publication No. US20040018545A1 GENERAL INFORMATION: APPLICANT: GONG, FANGCheng et al. FITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC FITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS, AND USES FILE REFERENCE: CL001195DIV2 CURRENT APPLICATION NUMBER: US/10/622,516 CURRENT APPLICATION NUMBER: US/10/622,516 CURRENT APPLICATION NUMBER: 10/193,295 PRIOR APPLICATION NUMBER: 10/193,295 PRIOR FILING DATE: 2002-07-12 PRIOR FILING DATE: 2002-07-12 PRIOR FILING DATE: 2001-03-29 NUMBER OF SEQ ID NOS: 5 SOFTWARE: FASTSEQ for Windows Version 4.0 SEQ ID NO 1 LENGTH: 2002	Qy 1261 AGAACTGGTGTGGCACCAGATGTCTTTAAAACATGAAGCTCAGAGGACCCCAT 1320 1261 AGAACTGGTTGGCACCAGATGTCTTTTAAACATGAAGCTCAGAGGACGACCCCAT 1320 1271 CATTTGGTCAACTATATTCCCCAGGGTTCAATAGATCACCATTTGAAGGAAG

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Qy 1 Db 1	<u> </u>	OV 1		Db 1	Qy 1	QY 1	ь	0v b			рь 14 Оу 12		Δρ 13		Qy 1	Qy 10	Db 11	·	Qy Db 1	ш	·	· L		p Qy	ДУ
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Qy 194 TGGGAATTGTTGCCCTTGAGATCTATTTTCCTTCTCAATATGTTGATCAAGCAGAGTTGG 253	134 GCTCTTTCACCATGCCTGGATCACTTCCTTTGAATGCAGAAGCTTGCTGGCCAAAAGATG 193 	Query Match 68.4%; Score 1370; DB 13; Length 1650; Best Local Similarity 92.0%; Pred. No. 0; Matches 1509; Conservative 0; Mismatches 5; Indels 126; Gaps 1;	TURE: E/KEY: CDS ATION: (22)(1582 36-417-181	ID NO 181 ENGTH: 1650 YPE: DNA RGANISM: Homo	PRIOR FILING DATE: 2001-09-19 Remaining Prior Application data remo NUMBER OF SEQ ID NOS: 341 SOFTWARE: Custom	PRIOR FILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/322,81 PRIOR FILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/323,51	PRIOR FILING DATE: 2002-07-17 PRIOR APPLICATION NUMBER: US60/322,63 PRIOR FILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/322,63	IOR FILING DATE: 2001-09-07 IOR APPLICATION NUMBER: US60/361,66 IOR FILING DATE: 2002-03-05 IOR APPLICATION NUMBER: US60/396,41	PRIOR FILING DATE: 2001-09-10 PRIOR APPLICATION NUMBER: US60/322,7 PRIOR FILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/318.1	PRIOR APPLICATION NUMBER: UPRIOR FILING DATE: 2001-09- PRIOR APPLICATION NUMBER: UPRIOR APPLICATION NUMBER: UPPRIOR APPLICATION APPLICATIO	FILE REFERENCE: 21402-442C CURRENT APPLICATION NUMBER: US/10/236,417	APPLICANT: Agee et al. TITLE OF INVENTION: NOVE	Sequence 181, Application No. US2	3 6	1988 AAAAAA 1994	Qy 1928 TAATCTCCAATTAAAAATTTTTTAACATGTAAAAAAAAAA		1868 TTTGGTTTTTAAACATGGTATAATGAATTGTGTACTTCTGTCAGAAGAAAGCAGAGGTAC 1	Qy 1808 CGAATGATGTTAAGGGCTCTGTAAAACTTCATACCTCTTTGGCCATTTGTATGCATGATG 1867	QY 1748 TCTCTTTGCTCTATTTGCTGACATGCTTCCTGTTGTGGTCTGGCCAATGCCAAATGTACT 1807
Qy 1148 CTGGT Db 1151 CTGGT	1088 TAG	Oy 1028 TATCA Db 1031 TATCA	968 AGGC 971 AGGC	Qy 908 GCCTG	Qy 848 GGATG Db 851 GGATG	Qy 788 ATTTT	Qy 728 GCAAA Db 731 GCAAA	Qy 668 ATGGA Db 671 ATGGG	Qy 608 ATATG	υλ 551 TAGCT		Ωу 592	Qy 554 TCAAT	Qy 494 ATACA Db 371 ATACA	Db 311 TCATC	CAT	Qy 374 TGGAG		AAA AAA	ДЬ 71 TTGGG.
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Sequence 183, Applic publication No. US20 GENERAL INFORMATION:
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LE REFERENCE: 21402-442C
RRENT APPLICATION NUMBER: US/10/236,417
RRENT FILING DATE: 2003-01-06
IOR APPLICATION NUMBER: US60/318,120
IOR FILING DATE: 2001-09-01
IOR APPLICATION NUMBER: US60/318,430
IOR FILING DATE: 2001-09-10
IOR APPLICATION NUMBER: US60/322,781
IOR FILING DATE: 2001-09-17
IOR APPLICATION NUMBER: US60/361,663
IOR APPLICATION NUMBER: US60/361,663
IOR APPLICATION NUMBER: US60/396,412
IOR APPLICATION NUMBER: US60/396,412
IOR APPLICATION NUMBER: US60/322,636
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AGANT: Agee et al.

E OF INVENTION: NOVEL PROTEINS AS REFERENCE: 21402-442C

LENT APPLICATION NUMBER: US/10/23

LENT FILING DATE: 2003-01-06

PRAPPLICATION NUMBER: US60/318,1

PRAPPLICATION NUMBER: US60/318,1

PRAPPLICATION NUMBER: US60/318,4

PRAPPLICATION NUMBER: US60/322,7

PRAPPLICATION NUMBER: US60/322,7

PRAPPLICATION NUMBER: US60/322,7

PRAPPLICATION NUMBER: US60/321,6

PRAPPLICATION NUMBER: US60/361,6

PRAPPLICATION NUMBER: US60/361,6

PRAPPLICATION NUMBER: US60/396,4

PRAPPLICATION NUMBER: US60/322,6

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; Sequence 189, Application US/10236417

; Publication No. US20040048256A1

; Publication No. US20040048256A1

; GENERAL INFORMATION:

; APPLICANT: Agee et al.

; TITLE OF INVENTION: NOVEL PROTEINS AND NU; FILE REFERENCE: 21402-442C

; CURRENT APPLICATION NUMBER: US/10/236,417

; PRIOR APPLICATION NUMBER: US60/318,120

; PRIOR FILING DATE: 2001-09-01

; PRIOR APPLICATION NUMBER: US60/318,430
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PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/318,184
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PRIOR FILING DATE: 2001-09-07
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR FILING DATE: 2002-03-05
PRIOR FILING DATE: 2002-07-17
PRIOR APPLICATION NUMBER: US60/396,412
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,817
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PRIOR APPLICATION NUMBER: US60/322,816
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/323,519
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PRIOR APPLICATION NUMBER: US60/323,519
PRIOR FILING DATE: 2001-09-19
Remaining Prior Application data removed - NUMBER OF SEQ ID NOS: 341
SOFTWARE: Custom
SEQ ID NO 189
LENGTH: 1650
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: (22)..(1582)
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US-10-236-417-193

US-10-236-417-193

Sequence 193, Application US/10236417

Publication No. US20040048256A1

GENERAL INFORMATION:
APPLICANT: Agee et al.

TITLE OF INVENTION: NOVEL PROTEINS AND NUTITLE OF INVENTION: NUMBER: US60/318,120

PRIOR APPLICATION NUMBER: US60/318,120

PRIOR APPLICATION NUMBER: US60/318,120

PRIOR APPLICATION NUMBER: US60/318,430

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR APPLICATION NUMBER: US60/361,663

PRIOR APPLICATION NUMBER: US60/396,412

PRIOR APPLICATION NUMBER: US60/322,636

PRIOR APPLICATION NUMBER: US60/322,636

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60/323,519

PRIOR FILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/323,519

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60
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US-10-236-417-197

i Sequence 197, Application US/10236417

i Publication No. US20040048256A1

i GENERAL INFORMATION:

APPLICANT: Agee et al.

i TITLE OF INVENTION: NOVEL PROTEINS AND NUCLEIC ACI

FILE REFERENCE: 21402-442C

CURRENT APPLICATION NUMBER: US/10/236,417

CURRENT PILING DATE: 2003-01-06

PRIOR APPLICATION NUMBER: US60/318,430

PRIOR FILING DATE: 2001-09-01

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR PILING DATE: 2001-09-10

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR RILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR RILING DATE: 2001-09-07

PRIOR APPLICATION NUMBER: US60/322,636

PRIOR PILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/396,412

PRIOR APPLICATION NUMBER: US60/322,636

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60/322,816

PRIOR FILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/322,816

PRIOR PILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/322,816

PRIOR PILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/323,519

PRIOR PILING DATE: 2001-09-19

Remaining PRIOR APPLICATION NUMBER: US60/323,519

PRIOR PILING DATE: 2001-09-19

REMAINING PRIOR APPLICATION NUMBER: US60/323,519

PRIOR PILING DATE: 2001-09-19

REMAINING PRIOR APPLICATION NUMBER: US60/323,519

PRIOR PILING DATE: 2001-09-19

REMAINING PRIOR APPLICATION NUMBER: US60/323,519

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RENT FILING DATE: 2003-01-06
R APPLICATION NUMBER: US60/318,120
R FILING DATE: 2001-09-01
R APPLICATION NUMBER: US60/318,430
R FILING DATE: 2001-09-10
R APPLICATE: 2001-09-10
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DR FILING DATE: 2002-03-05

DR APPLICATION NUMBER: US60/396,412

DR FILING DATE: 2002-07-17

DR APPLICATION NUMBER: US60/322,636

DR FILING DATE: 2001-09-17

DR APPLICATION NUMBER: US60/322,817

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RESULT 11 US-10-236-417-205 (Sequence 205, Application US/10236417 publication No. US20040048256A1 publication No. US20040048256A1 GENERAL INFORMATION: APPLICANT AGREE 201 FILE REFERENCE: 21402-442C CURRENT APPLICATION NOWBER: US/10/236,417 CURRENT FILING DATE: 2003-01-06 PRIOR PELICATION UNMBER: US60/318,430 PRIOR APPLICATION UNMBER: US60/318,430 PRIOR PILING DATE: 2001-09-10 PRIOR APPLICATION NUMBER: US60/318,430 PRIOR PILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/318,184 PRIOR PILING DATE: 2001-09-07 PRIOR APPLICATION NUMBER: US60/318,184 PRIOR PILING DATE: 2002-03-05 PRIOR APPLICATION NUMBER: US60/318,184 PRIOR APPLICATION NUMBER: US60/381,663 PRIOR PILING DATE: 2002-03-05 PRIOR APPLICATION NUMBER: US60/396,412 PRIOR APPLICATION NUMBER: US60/396,412 PRIOR APPLICATION NUMBER: US60/322,536 PRIOR APPLICATION NUMBER: US60/322,817 PRIOR APPLICATION NUMBER: US60/322,817 PRIOR APPLICATION NUMBER: US60/322,816 PRIOR PRILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/322,816 PRIOR PRILING DATE: 2001-09-17 PRIOR PRICE PRICE PRICE PRIOR PRICE PRI	Db 1091 TAGGACAGTACTCACCTCAGCAATTAGCAGGAAGAGAATTGGAGTGTTTCTTATGGTT 1150 Qy 1148 CTGGTTTGGCTGCCACTCTGTACTCTTTAAAGTCACAAGATGCTACACCAGGGTTCT 1207 Db 1151 CTGGTTTGGCTGCCACTCTGTACTCACTAAGTCACAAGATGCTACACCAGGGTTCT 1210 Qy 1208 CTCTTGATAAATAACAGCAAGTTTATGTGATCTTAAATCAAGACTGTACACCAGGGTTCT 1210 Qy 1208 CTCTTGATAAATAACAGCAAGTTTATGTGATCTTAAATCAAGACTTGATTCAAGAACTG 1267 Db 1211 CTCTTGATAAATAACAGCAAGTTTATGTGATCTTAAATCAAGGCTTGATTCAAGAACTG 1270 Qy 1268 GTGTGGCACCAGATGTCTTGCCTGAAAACATGAAGGCTCAGAAGAGCACCCATCATTTGG 1227 Qy 1276 GTGTGGCACCAGATGTCTCTCGCTGAAAACATGAAGCTCAGAAGAGACACCCATCATTTGG 1227 Qy 1287 TCAACTATATCCCCCAGGGTTCAATACATGAAGCTCAGAAGAGAACACCCATCATTTGG 1239 Db 1331 TCAACTATATTCCCCCAGGGTTCAATACATTCACTCTTTGAAGGAACACCCATCATTTGG 1330 Qy 1388 GGGTGGATGAAAACCACAGGAAACTTCACTCTTTGAAGGAACACGTGGATACTTACT

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RESULT 12
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; Publication No. US20040058338A1
; GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND FILE REFERENCE: 21402-502C
CURRENT APPLICATION NUMBER: US/10/307,
CURRENT FILING DATE: 2002-12-02
; NUMBER OF SEQ ID NOS: 682
; SOFTWARE: CuraSeqList version 0.1
; SEQ ID NO 359
LENGTH: 1650
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: (22)..(1581)
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US-10-307-817-371
; Sequence 371, Application US/10307817
; Publication No. US20040058338A1
; GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND NUCLE
FILE REFERENCE: 21402-502C
CURRENT APPLICATION NUMBER: US/10/307,817
; CURRENT FILING DATE: 2002-12-02
; NUMBER OF SEQ ID NOS: 682
; SOFTWARE: CuraSeqList version 0.1
; SEQ ID NO 371
; LENGTH: 1650
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
; LOCATION: (22)..(1581)
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US-10-307-817-445
; Sequence 445, Application US/10307817
; Publication No. US20040058338A1
; GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND NUTION OF ILE REFERENCE: 21402-502C
CURRENT APPLICATION NUMBER: US/10/307,817
CURRENT FILING DATE: 2002-12-02
; NUMBER OF SEQ ID NOS: 682
; SOFTWARE: CuraSeqList version 0.1
; SEQ ID NO 445
LENGTH: 1650
TYPE: DNA
; ORGANISM: Homo sapiens
US-10-307-817-445
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Sequence 372, Ap	Sequence 370, App	Sequence 360, App	Sequence 208, App	nence 206,	Sequence 204, App	uence 202, Ap	Sequence 198, App	Sequence 194, App	Sequence 190, App	Sequence 186, App	Sequence 184, Ap	Sequence 182, App	Sequence 2, Appl	, App	Description

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ce 284,	equence 316,	Sequence 208, App	ce 174,	equence 296,	nce 315,	e 314,	e 313,	e 14, A	equence 5,	ce 5, Ap	nce 36	e 358,	ce 210,	equence 200,	ce 376,	equence 366,	ce 212,	equence 196,	equence 362,	ce 188,	ce 364,	nce 192,	Ce	Sequence 56, Appl	, Ap	nce 292,	nce 468, Ap	ence 467,	Sequence 374, App

ALIGNMENTS

RESULT 1

US-10-193-295-2

US-10-193-295-2

; Sequence 2, Application US/10193295
; Publication No. US20020173018A1

; GENERAL INFORMATION:

; APPLICANT: GONG, Fangcheng et al.

; TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC

TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CLO01195DIV

; CURRENT APPLICATION NUMBER: US/10/193,295

; CURRENT APPLICATION NUMBER: 08/819,993
; PRIOR APPLICATION NUMBER: 08/819,993
; PRIOR FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 478
; TYPE: PRT
; ORGANISM: Human
US-10-193-295-2 \$ **D** S Вb Ş Query Match 100.0%; Best Local Similarity 100.0%; Matches 478; Conservative 0 121 121 61 61 \vdash MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCT o ; Score 2511; DB 13; Pred. No. 5.2e-241; Mismatches 0; Indels Length 478; o ; Gaps AND USES 180 180 120 120 60 09 0

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RESULT 2
US-10-622-516-2
US-10-622-516-2
; Sequence 2, Application US/10622516
; Publication No. US20040018545A1
; GENERAL INFORMATION: US20040018545A1
; GENERAL INFORMATION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEILLE OF INVENTION: THEREOF FILE REFERENCE: CL001195DIV2
; CURRENT APPLICATION NUMBER: US/10/622,516
; CURRENT FILING DATE: 2003-07-21
; PRIOR APPLICATION NUMBER: 10/193,295
; PRIOR FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: 09/819,993
; PRIOR FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 478
; TYPE: PRT
; ORGANISM: Human
US-10-622-516-2
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NMYTSSVYGSLASVLAQYSPQQLAGKRIGVFSYGSGLAATLYSLKVTQDATPGSALDKIT
                                                                          EGIDTTNACYGGTAAVFNAVNWIESSSWDGLRGTHMQHAYDFYKPDMLSEYPIVDGKLS
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                                    ALDRCYSVYCKKIHAQWQKEGNDKDFTLNDFGFMIFHSPYCKLVQKSLARMLLND
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US-10-236-417-182

US-10-236-417-182

; Sequence 182, Application US/10236417
; Publication No. US20040048256A1
; GENERAL INFORMATION:

TITLE OF INVENTION: NOVEL PROTEINS AND NUCLE
FILE REFERENCE: 21402-442C
CURRENT FILING DATE: 2003-01-06
PRIOR APPLICATION NUMBER: US60/318,120
PRIOR FILING DATE: 2001-09-01
PRIOR FILING DATE: 2001-09-10
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR FILING DATE: 2001-09-07
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR APPLICATION NUMBER: US60/326,412
PRIOR APPLICATION NUMBER: US60/396,412
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/323,519
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APPLICANT: Agee et al TITLE OF INVENTION: 1
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Matches 478
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TYPE: PRT
ORGANISM:
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RENT APPLICATION NUMBER: US/10/236,417
RENT FILING DATE: 2003-01-06
OR APPLICATION NUMBER: US60/318,120
OR APPLICATION NUMBER: US60/318,430
OR APPLICATION NUMBER: US60/318,430
OR APPLICATION NUMBER: US60/322,781
OR APPLICATION NUMBER: US60/322,781
OR APPLICATION NUMBER: US60/322,781
OR APPLICATION NUMBER: US60/318,184
OR APPLICATION NUMBER: US60/318,184
OR APPLICATION NUMBER: US60/318,184
OR APPLICATION NUMBER: US60/318,184
OR APPLICATION NUMBER: US60/322,636
OR APPLICATION NUMBER: US60/322,636
OR APPLICATION NUMBER: US60/322,636
OR APPLICATION NUMBER: US60/322,817
OR APPLICATION NUMBER: US60/322,817
OR APPLICATION NUMBER: US60/322,816
OR FILING DATE: 2001-09-17
OR APPLICATION NUMBER: US60/323,519
OR APPLICATION NUMBER: US60/323,519
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o. US20040048256A1
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Application
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CURRENT FILING DATE: 2003-01-06
PRIOR APPLICATION NUMBER: US60/318,120
PRIOR FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR FILING DATE: 2001-09-10
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR FILING DATE: 2001-09-07
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR APPLICATION NUMBER: US60/396,412
PRIOR APPLICATION NUMBER: US60/396,412
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/323,519
                                                                                                                                                    SOFTWARE:
SEQ ID NO 1
LENGTH: 5
TYPE: PRT
ORGANISM:
S-10-236-417
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                                                                                                                                                       NISM: Homo 6-417-186
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                                                                                                                                                                                                                                 FILING DATE: 2001-09-19
ing Prior Application data
OF SEQ ID NOS: 341
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DREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDI
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RESULT 6
US-10-236-417-190

Sequence 190, Application US/10236417

Publication No. US20040048256A1

GENERAL INFORMATION:
APPLICANT: Agee et al.
FILLE REFERENCE: 21402-442C
CURRENT APPLICATION NOVEL PROTEINS AND NUMBER: US/10/236,417
CURRENT FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US/0/318,120
PRIOR APPLICATION NUMBER: US/0/318,430
PRIOR FILING DATE: 2001-09-01
PRIOR FILING DATE: 2001-09-10
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US/0/322,781
PRIOR APPLICATION NUMBER: US/0/322,781
PRIOR APPLICATION NUMBER: US/0/318,184
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US/0/361,663
PRIOR FILING DATE: 2002-03-05
PRIOR APPLICATION NUMBER: US/0/326,412
PRIOR APPLICATION NUMBER: US/0/322,636
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US/0/322,817
PRIOR APPLICATION NUMBER: US/0/322,816
PRIOR FILING DATE: 2001-09-17
                       Query Match

Best Local Similarity 91.

Matches 478; Conservative
                                                                                                           LENGTH: 520
TYPE: PRT
ORGANISM: Homo
10-236-417-190
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US-10-236-417-194
; Sequence 194, Application US/10236417
; Publication No. US20040048256A1
; GENERAL INFORMATION:
APPLICANT: Agee et al.
FITTLE OF INVENTION: NOVEL PROTEINS AND NUCLE
FILE REFERENCE: 21402-442C
CURRENT FILING DATE: 2003-01-06
PRIOR APPLICATION NUMBER: US/10/236,417
CURRENT FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR FILING DATE: 2001-09-10
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR FILING DATE: 2002-03-05
PRIOR APPLICATION NUMBER: US60/364,12
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR APPLICATION NUMBER: US60/322,817
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PRIOR APPLICATION NUMBER: US60/322,816
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION PUMBER: US60/323,519
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Sequence 198, Application US/10236417; Publication No. US20040048256A1; GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND NUFILE REFERENCE: 21402-442C
CURRENT APPLICATION NUMBER: US/10/236,417
CURRENT FILING DATE: 2003-01-06
PRIOR APPLICATION NUMBER: US60/318,120
PRIOR FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR FILING DATE: 2001-09-10
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR FILING DATE: 2001-09-07
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,636
PRIOR APPLICATION NUMBER: US60/322,636
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NUMBER OF SEQ |
SOFTWARE: Custor |
SEQ ID NO 198 |
LENGTH: 520 |
TYPE: PRT |
ORGANISM: Houseld |
S-10-236-417-19
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US-10-236-417-202
US-10-236-417-202
; Sequence 202, Application US/10236417
; Publication No. US20040048256A1
; GENERAL INFORMATION:
APPLICANT: Agee et al.
; TITLE OF INVENTION: NOVEL PROTEINS AND NUCLEIC ACIDS ENCODING SAPILE REFERENCE: 21402-442C
; CURRENT APPLICATION NUMBER: US/10/236,417
; CURRENT FILING DATE: 2003-01-06
; PRIOR APPLICATION NUMBER: US60/318,120
; PRIOR APPLICATION NUMBER: US60/318,430
; PRIOR APPLICATION NUMBER: US60/318,430
; PRIOR APPLICATION NUMBER: US60/322,781
; PRIOR APPLICATION NUMBER: US60/326,412
; PRIOR APPLICATION NUMBER: US60/361,663
; PRIOR APPLICATION NUMBER: US60/361,663
; PRIOR APPLICATION NUMBER: US60/364,412
; PRIOR FILING DATE: 2002-03-05
; PRIOR FILING DATE: 2002-07-17
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GENERAL INFORMATION:

APPLICANT: Agee et al.

TITLE OF INVENTION: NOVEL PROTEINS AND NI

FILE REFERENCE: 21402-442C

CURRENT APPLICATION NUMBER: US/10/236,417

CURRENT FILING DATE: 2003-01-06

PRIOR APPLICATION NUMBER: US60/318,120

PRIOR APPLICATION NUMBER: US60/318,430

PRIOR APPLICATION NUMBER: US60/322,781

PRIOR APPLICATION NUMBER: US60/322,781
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R FILING DATE: 2001-09-17
R APPLICATION NUMBER: US60/322,817
R FILING DATE: 2001-09-17
R APPLICATION NUMBER: US60/322,816
R APPLICATION NUMBER: US60/322,816
R FILING DATE: 2001-09-17
R APPLICATION NUMBER: US60/323,519
R FILING DATE: 2001-09-19
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PRIOR FILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/318,184

PRIOR APPLICATION NUMBER: US60/361,663

PRIOR FILING DATE: 2002-03-05

PRIOR APPLICATION NUMBER: US60/361,663

PRIOR FILING DATE: 2002-07-17

PRIOR APPLICATION NUMBER: US60/322,636

PRIOR FILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60/322,817

PRIOR APPLICATION NUMBER: US60/322,816

PRIOR APPLICATION NUMBER: US60/323,519

PRIOR APPLICATION NUMBER: US60/323,519

PRIOR FILING DATE: 2001-09-17

PRIOR FILING DATE: 2001-09-19

Remaining Prior Application data removed NUMBER OF SEQ ID NOS: 341

SOFTWARE: Custom

SOFTWARE: Custom

TYPE: PRI
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RESULT 11
US-10-236-417-206
; Sequence 206, Application US/
; Publication No. US2004004825(
; GENERAL INFORMATION:
; APPLICANT: Agee et al.
; TITLE OF INVENTION: NOVEL P.
; FILE REFERENCE: 21402-442C

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CURRENT FILING DATE: 2003-01-06
PRIOR APPLICATION NUMBER: US60/318,120
PRIOR FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR FILING DATE: 2001-09-10
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/318,184
PRIOR FILING DATE: 2001-09-07
PRIOR FILING DATE: 2001-09-07
PRIOR FILING DATE: 2002-03-05
PRIOR FILING DATE: 2002-07-17
PRIOR APPLICATION NUMBER: US60/396,412
PRIOR FILING DATE: 2002-07-17
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,817
PRIOR APPLICATION NUMBER: US60/322,816
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR APPLICATION NUMBER: US60/323,519
PRIOR FILING DATE: 2001-09-17
PRIOR FILING DATE: 2001-09-17
PRIOR PILING DATE: 2001-09-19
Remaining Prior Application data remove NUMBER OF SEQ ID NOS: 341
SOFTWARE: Custom
SEQ ID NOS: 341
ORGANISM: Homo sapiens
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RRENT FILING DATE: 2003-01-06
IOR APPLICATION NUMBER: US60/318,120
IOR APPLICATION NUMBER: US60/318,430
IOR APPLICATION NUMBER: US60/318,430
IOR FILING DATE: 2001-09-10
IOR APPLICATION NUMBER: US60/322,781
IOR FILING DATE: 2001-09-17
IOR APPLICATION NUMBER: US60/322,781
IOR APPLICATION NUMBER: US60/318,184
IOR FILING DATE: 2001-09-07
IOR APPLICATION NUMBER: US60/361,663
IOR FILING DATE: 2002-03-05
IOR FILING DATE: 2002-07-17
IOR APPLICATION NUMBER: US60/322,636
IOR APPLICATION NUMBER: US60/322,817
IOR APPLICATION NUMBER: US60/322,817
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IOR APPLICATION NUMBER: US60/323,519
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IOR SEQ ID NOS: 341
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RESULT 13
US-10-307-817-360
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; GENERAL INFORMATION:
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TITLE OF INVENTION: NOVEL PRO
FILE REFERENCE: 21402-502C
CURRENT APPLICATION NUMBER: U
CURRENT FILING DATE: 2002-12
NUMBER OF SEQ ID NOS: 682
SOFTWARE: CuraSeqList version
SEQ ID NO 360
LENGTH: 520
                           Sequence 370, Applic Publication No. US20 GENERAL INFORMATION:
APPLICANT: Agee et al. TITLE OF INVENTION: NO FILE REFERENCE: 21402-
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              0-307-817-
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ORGANISM: Homo
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CURRENT APPLICATION NUMBER: US/10/307,8:
CURRENT FILING DATE: 2002-12-02
NUMBER OF SEQ ID NOS: 682
SOFTWARE: CuraSeqList version 0.1
SEQ ID NO 370
LENGTH: 520
TYPE: PRT
ORGANISM: Homo sapiens
US-10-307-817-370
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                                             Sequence 372, Application US/10307817; Publication No. US20040058338A1; GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND NOTE OF SEQION NUMBER: US/10/307,81; CURRENT APPLICATION NUMBER: US/10/307,81; CURRENT FILING DATE: 2002-12-02; NUMBER OF SEQID NOS: 682; SOFTWARE: CuraSeqList version 0.1; SEQID NO 372; LENGTH: 520
TYPE: PRT
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Search completed: June 24, 2004, 13:44:07 Job time: 884 secs

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Sequence 1, Appli	Sequence 1, Appli	Sequence 13, Appl	, Appl	•	, Appl	, Appl	Sequence 5, Appli	, Appl	סי	, Appl	e 9, Appl	0, App	0, App	57,	, 98	Sequence 4, Appli	equence

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patent No. 6436692

GENERAL INFORMATION:
APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYME P
TITLE OF INVENTION: ACID MOLECULES ENCODING
TITLE OF INVENTION: THEREOF
FILE REFERENCE: CL001195
CURRENT APPLICATION NUMBER: US/09/819,993
CURRENT FILING DATE: 2001-03-29
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSEQ for Windows Version 4 ^
SEQ ID NO 2
LENGTH: 478
TYPE: PRT
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; Sequence 2, Application US/10193295
; Patent No. 6620608
; GENERAL INFORMATION:
APPLICANT: GONG, Fangcheng et al.
APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC
TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTE
TITLE OF INVENTION: THEREOF
FILE REFERENCE: CL001195DIV
CURRENT APPLICATION NUMBER: US/10/193,295
CURRENT FILING DATE: 2002-07-12
PRIOR APPLICATION NUMBER: 08/819,993
PRIOR FILING DATE: 2001-03-29

NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 2
LENGTH: 478
TYPE: PRT
ORGANISM: Human
US-10-193-295-2
 RESULT 3
US-09-819-993-4
; Sequence 4, Application US/0981
; Patent No. 6436692
; GENERAL INFORMATION:
; APPLICANT: GONG, Fangcheng et
; TITLE OF INVENTION: ISOLATED H
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Pred. No. 1.;
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RESULT 4
US-10-193-295-4
; Sequence 4, Application US/10193295
; Patent No. 6620608
; GENERAL INFORMATION:
APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NU
TITLE OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZ
TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001195DIV
; CURRENT APPLICATION NUMBER: US/10/193,295
; CURRENT FILING DATE: 2002-07-12
; PRIOR APPLICATION NUMBER: 08/819,993
; PRIOR FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 4
; LENGTH: 520
; TYPE: PRT
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TITLE OF INVENTION: ACID MOLECULES ENCO.
FILE REFERENCE: CL001195
CURRENT APPLICATION NUMBER: US/09/819,99:
CURRENT FILING DATE: 2001-03-29
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSEQ for Windows Version 4.1
SEQ ID NO 4
LENGTH: 520
TYPE: PRT
ORGANISM: Human
US-09-819-993-4
                                                                                                                                                                                                                                                                                                                      98.8%; l Similarity 91.9%; 478; Conservative
VGLVHSNIATEHIPSPAKKVPRLPATAAEPEAAVISNGEH 478
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Pred. No. 3.1e-242;
); Mismatches 0;
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WS-09-819-993-5
; Sequence 5, Application US/09819993
; Patent No. 6436692
; GENERAL INFORMATION:
APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZ
TITLE OF INVENTION: ACID MOLECULES EN
TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001195
; CURRENT APPLICATION NUMBER: US/09/819,
; CURRENT FILING DATE: 2001-03-29
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version
; SEQ ID NO 5
; LENGTH: 518
; TYPE: PRT
; ORGANISM: Human
US-09-819-993-5
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JAPLICANT: GONG, Fangcheng et al.

JAPLICANT: GONG, Fangcheng et al.

JITLE OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUTTITLE OF INVENTION: THEREOF

FILE REFERENCE: CLOO1195DIV

CURRENT APPLICATION NUMBER: US/10/193,295

CURRENT APPLICATION NUMBER: 08/819,993

PRIOR APPLICATION NUMBER: 08/819,993

PRIOR PILING DATE: 2001-03-29

NUMBER OF SEQ ID NOS: 5

SOFTWARE: FastSEQ for Windows Versing Corganism: 518

TYPE: PRT

ORGANISM: Himm

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Similarity 90.9%;
71; Conservative
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                                  Score 2439; DB 4;
Pred. No. 4.3e-238;
1; Mismatches 4;
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                                                                                                                                                                                                                                                                                                                                TELECOMMUNICATION INFORMATION:
TELEPHONE: (414) 277-5709
TELEFAX: (414) 277-5591
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 520 amino acids
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Patent No
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GENERAL INFORMATION:
APPLICANT: Miziork
TITLE OF TANK
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CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/072,040
FILING DATE: 02 JUNE 1993
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 65-053-908:
TELECOMMUNICATION INFORMATION:
TELEPHONE: (414) 277-5709
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MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0,

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/305,505
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ADDRESSEE: Quarles & I
STREET: 411 East Wisco
CITY: Milwaukee
STATE: Wisco
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3-HYDROXY-3-METHYLGLUTARYL-COA
SYNTHASE PREPARATION WITH IMPROVED
STABILITY
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Pred. No. 4.4e-238;
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RESULT 8
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GENERAL INFORMATION:
GENERAL INFORMATION:
GENERAL INFORMATION:
APPLICANT: Miziorko, Henry M.
TITLE OF INVENTION: 3-HYDROXY-3-METH
TITLE OF INVENTION: SYNTHASE PREPARA
TITLE OF INVENTION: STABILITY
NUMBER OF SEQUENCES: 6
CORRESPONDENCE ADDRESS:
ADDRESSEE: Quarles & Brady
STREET: 411 East Wisconsin Avenue
CITY: Milwaukee
STATE: Wisconsin
COUNTRY: U.S.A.
ZIP: 53202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, V
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/305,505
FILING DATE:
CLASSIFICATION DATA:
APPLICATION NUMBER: US/08/072,040
FILING DATE: 02 JUNE 1993
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 65-053-90
TELECOMMUNICATION INFORMATION:
TELEPHONE: (414) 277-5591
INFORMATION FOR SEQ ID NO: 5:
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3-HYDROXY-3-METHYLGLUTARYL-COA
SYNTHASE PREPARATION WITH IMPROVED
STABILITY
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                            RESULT 9

IS-08-305-505-4

Sequence 4, Application US/08305505

Patent No. 5668001

GENERAL INFORMATION:
APPLICANT: Miziorko, Henry M.
TITLE OF INVENTION: 3-HYDROXY-3

TITLE OF INVENTION: SYNTHASE PRI
TITLE OF INVENTION: STABILITY

NUMBER OF SEQUENCES: 6

CORRESPONDENCE ADDRESS:
ADDRESSEE: Quarles & Brady

STREET: 411 East Wisconsin Av.
CITY: Milwaukee

STATE: Wisconsin

COUNTRY: U.S.A.
ZIP: 53202

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-D
SOFTWARE: PatentIn Release #1

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/305
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TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
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RESULT 10
US-08-305-505
; Sequence 2, App.;
; Patent No. 5668001
; Patent No. 5668001
; GENERAL INFORMATION:
; APPLICANT: Miziorko, He
TITLE OF INVENTION: 3-/
TITLE OF INVENTION: SY
; TITLE OF INVENTION: SY
; NUMBER OF SEQUENCES:
; CORRESPONDENCE ADDRES:
ADDRESSEE: Quarles
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CLAC

PRIOR APPL

APPLICATION

FILING DATE:

ATTORNEY/AGENT INFORM.

NAME: Baker, Jean C.

REGISTRATION NUMBER: 35,%

REFERENCE/DOCKET NUMBER: 65-C

TELECOMMUNICATION INFORMATION:

TELEPHONE: (414) 277-5709

TELEFAX: (414) 277-5709

TELEFAX: (414) 277-5591

INFORMATION FOR SEQ ID NO: 4:

SEQUENCE CHARACTERISTICS:

LENGTH: 520 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: protein

"NS-08-305-505-4"

Match

Match

Similarity 87.5%;

Conservative

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Pred. No. 2.8e
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Ce 2, Application US/08305505 No. 5668001 AL INFORMATION:

Henry M.
3-HYDROXY-3-METHYLGLUTARYLSYNTHASE PREPARATION WITH I
STABILITY

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CITY: Milwaukee
STATE: Wisconsin
COUNTRY: U.S.A.
ZIP: 53202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PATENTION PC-DOS/MS-DOS
SOFTWARE: PATENTION DATA:
APPLICATION NUMBER: US/08/305,505
FILING DATE:
CLASSIFICATION UMBER: US/08/072,040
FILING DATE: 02 JUNE 1993
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 65-053-908
TELECOMMUNICATION INFORMATION:
TELEPHONE: (414) 277-5591
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 522 amino acids
TYPE: amino acid
STRANDEDNESS: Single
TOPOLOGY: linear
MOLECULE TYPE: protein
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Pred. No. 2e-2
6; Mismatches
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RESULT 11
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; Patent No. 566
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ADDRESSEE: Quarles & Brady
STREET: 411 East Wisconsin Avenue
CITY: Milwaukee
STATE: Wisconsin
COUNTRY: U.S.A.
ZIP: 53202
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Ve
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/305,505
FILING DATE:
CLASSIFICATION DATA:
APPLICATION NUMBER: US/08/072,040
FILING DATE: 02 JUNE 1993
ATTORNEY/AGENT INFORMATION:
NAME: Baker, Jean C.
REGISTRATION NUMBER: 35,433
REFERENCE/DOCKET NUMBER: 65-053-908
TELEPHONE: (414) 277-5591
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 507 amino acids
TYPE: amino acids
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TITLE OF INVENTION:
TITLE OF INVENTION:
NUMBER OF SEQUENCES:
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the Local Similarity 58.9%; Proches 274; Conservative 76;
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STRANDEDNESS: sin
TOPOLOGY: linear
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No. 5668001
NAL INFORMATION:
PLICANT: Miziorko,
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3-HYDROXY-3-METHYLGLUTARYL-COA
SYNTHASE PREPARATION WITH IMPROVED
STABILITY
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Pred. No. 1.4e-140;
5; Mismatches 72;
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RESULT 13
US-09-925-388-6
; Sequence 6, F
; Patent No. 65
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APPLICANT: SETOGUCHI, Yutaka
TITLE OF INVENTION: ISOPRENOID PRODUCTION
FILE REFERENCE: ISOPRENOID PRODUCTION
CURRENT APPLICATION NUMBER: US/09/306,595C
CURRENT FILING DATE: 1999-05-06
PRIOR APPLICATION NUMBER: 98108210
PRIOR FILING DATE: 1998-05-06
NUMBER OF SEQ ID NOS: 43
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 6
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US-09-306-5
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Best Local
Matches 19
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APPLICANT: HOSHINO, Tat
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TYPE: PRT
ORGANISM: Ph
09-306-595C-6
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CANT: SETOGUCHI, Yutaka
CANT: SETOGUCHI, Yutaka
OF INVENTION: ISOPRENOID PRODUCTION
REFERENCE: ISOPRENOID PRODUCTION
NT APPLICATION NUMBER: US/09/925,388
NT FILING DATE: 2001-08-09
                                CANT: Wang, Tongtong
OF INVENTION: COMPOUNDS
OF INVENTION: DIAGNOSIS
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       REFERENCE: 210121.471C2
NT APPLICATION NUMBER: US/09/401,064
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CORRESPONDENCE ADDRESS:
ADDRESSE: GENOME THERAPEUTICS CORPORA
STREET: 100 Beaver Street
CITY: Waltham
STATE: Massachusetts
COUNTRY: USA
ZIP: 02354

COMPUTER READABLE FORM:
MEDLIUM TYPE: CD/ROM ISO9660
COMPUTER: PC
OPERATING SYSTEM: <Unknown>
SOFTWARE: ASCII
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/107,532A
FILING DATE: 30-Jun-1998
PRIOR APPLICATION NUMBER: 60/085,598
FILING DATE: 14 May 1998
APPLICATION NUMBER: 60/085,598
FILING DATE: 14 May 1998
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WPI N-P Gon 9Sn Human; enzyme; chromosome 5; hydroxymethylglutaryl-coenzyme A synthase; HMG-CoA synthase; cholesterologenesis; therapeutic; diagnostic; genotypantibody; synthase; carcinoma. T 1 726 ABG32726 standard; 29-(AP 20 Homo sapiens 22-Human hydroxymethylglutaryl-coenzyme ABG32726; 9 SDB; PL-) AUG-2002. 436692-B1. MAR-2001; MAR-2001; NOV-2002 , L) 2002-689940/74. B; ABS54409. APPLERA CORP. Yan C, 2001US-00819993. 2001US-00819993. (first entry) Di Francesco protein; 478 **,** AA. Beasley A synthase protein. genotype;

The invention discloses an isolated nucleic acid molecule encoding a human hydroxymethylglutaryl-coenzyme A (HMG-CoA) synthase which is important in cholesterologenesis. The polynucleotide and polypeptide are useful as models for the development of human therapeutic targets, to aid in the identification of therapeutic proteins and as targets for the development of human therapeutic agents that modulate the activity of the polypeptide in cells and tissues. The polynucleotide is useful for activity of the effectiveness of modulating compounds on the expression or activity of the enzyme gene in clinical trials and in treatments, in diagnostic assays for qualitative changes in expression of enzyme nucleic acid, to detect mutations in enzyme genes and gene expression products,

New syn

isolated nucleic acid molecule encoding hydroxymethylglutaryl-CoAthase, useful as model for the development of human therapeutic gets and for identifying therapeutic proteins.

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that shows a high degree of similarity to human cytoplasmic 3-hydroxy-3-methylglutaryl coenzyme A (FMG-CoA) synthase. The invention also discloses polynucleotide sequences encoding the novel enzyme of the invention. Both the polypeptide and polynucleotide sequences are useful as models for the development of human therapeutics, for identifying therapeutic proteins, as targets for development of human therapeutic as encoding the novel enzyme of the invention. Both the polypeptide and polynucleotide sequences are useful casematics, and as query sequences to perform a search against sequence data bases to identify other family members of related sequences. The polypeptide is useful to raise antibodies or to elicit another immune response, as a reagent in assays designed to quantitatively determine levels of the protein in biological fluids, as markers for tissues in which the corresponding protein is preferentially expressed, in drug state, or an altered form that causes the specific disease or pathology associated with the synthase activity of the protein in its natural conjugated with the synthase, to screen a compound for the ability to state, or an altered form that causes the specific disease or pathology associated with the synthase, to screen a compound for the ability to pharmacogenomic analysis. The polynucleotide is useful for monitoring the pharmacogenomic analysis. The polynucleotide is useful for monitoring the pharmacogenomic analysis. The polynucleotide is useful for activity of the human synthase gene in clinical trials or in a treatment regimen, in diagnostic assays for qualitative changes in a human synthase nucleic acid that leads to a pathology, for testing an individual for a genotype that while not necessarily causing a disease, nevertheless affects the creatment modality, and as antisense constructs to control human synthase gene in control human synthase gene captured the control human synthase gene antisense constructs to control human synthase.
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The present invention describe
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hypotensive, dermatological, a
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disorders, neoplasm, lymphoma,
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infectious disease, anorexia,
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ABR54277 represents a human to
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                                    antiparkinatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with the NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, valve disease, tuberous sclerosis, scleroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, AlDS, bronchial asthma, Crohn's disease, multiple sclerosis, infectious disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, Parkinson's disease, immune disorders, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention.

ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention
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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in CC ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, Dypotensive, dermatological, annorectic, immunosuppressive, cytostatic, antiidiabetic, antiinfertility, haemostatic, immunosuppressive, cytostatic, antiidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, CC antiparkinsonian and antilipaemic activities, and can be used in gene conting a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and can be used in the treatment and diagnosis of cardiomyopathy, anti-Hodies can be used in the treatment and diagnosis of cardiomyopathy, congenital adrenal hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, cc disease, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, parkinson's disease, immune disorders, cancer, alzheimer's disease, parkinson's disease, immune disorders, cancer, captaled acchexia, cancer, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, parkinson's disease, immune disorders, cancer, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, parkinson's disease, immune disorders, cancer, alzheimer's disease, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, al
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Ort T, Shimkets RA;
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J, Dipippo
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2001US-0323819P.
2001US-0323631P.
2001US-0323636P.
2001US-0324969P.
2001US-0324969P.
2001US-0324990P.
2001US-0341144P.
2002US-0361663P.
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    Boldog FL;
Chaudhuri A;
Ellerman K;
                      A,
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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in C ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antialdiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiatherosclerotic, antiinfertility, haemostatic, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital darenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, diabetes, metabolic disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, parkinson's disease, immune disorders, cancer, caccayate to ACC62465 represent PCR primers and metabolic syndrome X. ARS4277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemples from the present invention.
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Kekuda R, Khramtsov NV,
Malyankar UM, Miller CE,
Pena CEA, Rieger DK, Roti
Spaderna SK, Spytek KA,
Zerhusen BD, Zhong M;
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N-PSDB;
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Sequence
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B; ACC62339.
   520 AA;
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camtsov NV, Leach MD, Lepley DM, Li L, Liu X;
Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M;
ger DK, Rothenberg ME, Shenoy SG, Shimkets RA;
Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss E
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             247; 460pp; English.
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Query Match Best Local S Matches 478

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yankar UM, Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M;
la CEA, Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA;
derna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss Eihusen BD, Zhong M;
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HERERE KREET Gangolli EA, Gerlach VI, Giot I, Gol Kekuda R, Khramtsov NV, Leach MD, Le Malyankar UM, Miller CE, Ooi CE, Ort Pena CEA, Rieger DK, Rothenberg ME, Spaderna SK, Spytek KA, Taupier RJ, Zerhusen BD, Zhong M; Gorman L, Guo X, Gusev VY
Lepley DM, Li L, Liu X;
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E, Shenoy SG, Shimkets RA;
J, Twomlow N, Vernet CAM, Patturajan M; **5** Voss EZ; ďί Σ.

WPI; 200 N-PSDB; 2003-313241/30.)B; ACC62338.

Novel human proteins and nucleic acid encoding the proteins, useful for diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.

Claim 1; Page 246; 460pp; English.

The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, C hypotensive, dermatological, annorectic, immunosuppressive, cytostatic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, C antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, associated with a NOVX protein in humans and for treating a pathology c associated with the human disease. NOVX nucleic acids, proteins and atherosclerosis, hypertension, congenital heart defects, aortic stenosis, C valve disease, tuberous sclerosis, sclerosis, echeroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host c disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, hazheimer's disease, parkinson's disease, immune disorders, cancer, hazheimer's disease, parkinson's disease, immune disorders, haemophilia, chocalle to ACC62465 represent PCR primers and metabolic syndrome X. ACC62346 to ACC62455 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention. The human NOV35b protein in the exemplification of the present invention. Sequence 520 AA,

Query Match Best Local S Matches 478 Ch 98.8%; l Similarity 91.9%; 478; Conservative Score 2480; DB 6; Pred. No. 8.1e-234; 0; Mismatches 0; Length 520; Indels 42; Gaps

378	SPQQLAGKRIGVFSYGSGLAATLYSLKVTQDATPGSALDKITASLCDLKSRLDSRTGVAP :	319	γΩ
360	FGDVKLEDTYFDRDVEKAFMKASSELFSQKTKASLLVSNQNGNMYTSSVYGSLASVLAQY	301	Db
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300	HAQWQKEGNDKDFTLNDFGFMIFHSPYCKLVQKSLARMLLNDFLNDQNRDKNSIYSGLEA	241	В
258	HAQWQKEGNDKDFTLNDFGFMIFHSPYCKLVQKSLARMLLNDFLNDQNRDKNSIYSGLEA	199	VQ VQ
240	IGPNAPLIFERGLRGTHMQHAYDFYKPDMLSEYPIVDGKLSIQCYLSALDRCYSVYCKKI	181	Db
198	GLRGTHMQHAYDFYKPDMLSEYPIVDGKLSIQCYLSALDRCYSVYCKKI	150	Qy
180	EGIDTTNACYGGTAAVFNAVNWIESSSWDGRYALVVAGDIAVYATGNARPTGGVGAVALL	121	gg
149	EGIDTTNACYGGTAAVFNAVNWIESSSWD	121	Q
120	DREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDI	61	DЬ
120	DREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDI	61	γQ
60	MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCT	–	DЬ
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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, arctic stenosis, valve disease, tuberous sclerosis, sclerosis, scleroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, lacked, to ACC62346 to ACC62465 represent PCR primers and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX and the properties of the properties of the properties of the properties of the properties.
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CE, Ooi CE, Ort T, Padigaru M, Patturajan M;
Rothenberg ME, Shenoy SG, Shimkets RA;
A, Taupier RJ, Twomlow N, Vernet CAM, Voss E
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Gangolli EA, Kekuda R, Khr Malyankar UM, Pena CEA, Rie Spaderna SK, Zerhusen BD, EA, Gerlach VL, Giot L, Gorman L, Guo X, Gusev VY, Ji W, Khramtsov NV, Leach MD, Lepley DM, Li L, Liu X; r UM, Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M; Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA; SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss EBD, Zhong M; Voss EZ;

WPI; 200 N-PSDB; B; ACC62336.

Novel human proteins and nucleic acid encoding the proteins, useful diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.

Claim 1; Page 245; 460pp; English.

The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, C hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiinfertility, haemostatic, inmunosuppressive, cytostatic, antiinfertility, haemostatic, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, diabetes, metabolic hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host confectious disease, anorexia, cancer -associated cachexia, cancer, haemophilia, confectious disease, anorexia, cancer -associated cachexia, cancer, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62346 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention.

Charactery and the present invention of the present invention with the human NOV35b protein in the exemplification of the present invention. Sequence 520 AA;

Que Bes Mat ery Match 98.8%; Local Similarity 91.9%; ches 478; Conservative 301 199 181 241 121 61 -MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCT HAQWQKEGNDKDFTLNDFGFMIFHSPYCKLVQKSLARMLLNDFLNDQNRDKNSIYSGLEA FGDVKLEDTYFDRDVEKAFMKASSELFSQKTKASLLVSNQNGNMYTSSVYGSLASVLAQY DREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDI SPQQLAGKRIGVFSYGSGLAATLYSLKVTQDATPGSALDKITASLCDLKSRLDSRTGVAP ------GLRGTHMQHAYDFYKPDMLSEYPIVDGKLSIQCYLSALDRCYSVYCKKI EGIDTTNACYGGTAAVFNAVNWIESSSWD--------EGIDTTNACYGGTAAVFNAVNWIESSSWDGRYALVVAGDIAVYATGNARPTGGVGAVALL Score 2480; DB 6; Pred. No. 8.1e-234; 0; Mismatches 0; Indels 42; Gaps 198 300 258 240 180 60 360 120 149 120 09 <u>,</u>

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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antidabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atheroselerosis, hypertension, congenital heart defects, aortic stenosis, valve disease, tuberous sclerosis, sclerosers, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, diabetes, metabolic disease, allows, idiopathic thrombocytopenic purpura, graft versus host chiesease, AlDS, bronchial asthma, Crohn's disease, multiple sclerosis, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention. The human NOV35b protein in the exemplification of the present invention.
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Khramtsov NV, Lead...
UM, Miller CE, Ooi CE, Ort
Rieger DK, Rothenberg ME, Si
Rieger NA, Taupier RJ, T
  520 AA;
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Lepley DM,
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T, Padigaru M, Patturajan M;
Shenoy SG, Shimkets RA;
Twomlow N, Vernet CAM, Voss E
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S ঠ Ş 밁 S 멂 S 8 Ş 9 B 명 Query Match Best Local Matches 47 319 301 241 199 181 121 478; 61 \vdash Similarity MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCT HAQWQKEGNDKDFTLNDFGFMIFHSPYCKLVQKSLARMLLNDFLNDQNRDKNSIYSGLEA EGIDTTNACYGGTAAVFNAVNWIESSSWD--DREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDI DREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSKSVKTNLMQLFEESGNTDI HAQWQKEGNDKDFTLNDFGFM1FHSPYCKLVQKSLARMLLNDFLNDQNRDKNS1YSGLEA -----GLRGTHMQHAYDFYKPDMLSEYPIVDGKLSIQCYLSALDRCYSVYCKKI SPQQLAGKRIGVFSYGSGLAATLYSLKVTQDATPGSALDKITASLCDLKSRLDSRTGVAP Conservative 98.8%; 91.9%; Score 2480; DB 6; Pred. No. 8.1e-234; 0; Mismatches 0; Length 520; Indels 42; Gaps 120 360 300 240 198 180 149 60 60 258

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26-SEP-2002;

03-MAY-2002;

03-MAY-2002;

17-JUL-2002;

13-AUG-2002;

16-SEP-2002;
        Agee ML, Al
Burgess CE,
Crabtree J,
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2001US-0324969P.
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2001US-0341144P.
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Gangolli EA, Gerlach VL, Giot L, Gorman L, Guo X, Gusev VY, Ji W Kekuda R, Khramtsov NV, Leach MD, Lepley DM, Li L, Liu X; Malyankar UM, Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M; Pena CEA, Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA; Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss El Zerhusen BD, Zhong M;

WPI; 2003-313241/30. N-PSDB; ACC62334.

Novel human proteins and nucleic acid encoding the proteins, useful diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.

Page 244; 460pp; English.

HERETE KREET The present invention describes isolated human NOVX proteins, where X is CC 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in CC ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, CC hypotensive, dermatological, ancerctic, immunosuppressive, cytostatic, CC antidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, CC antisthmatic, metabolic, immunomodulator, neuroprotective, nootropic, CC antiparkinsonian and antilipaemic activities, and can be used in gene CC associated with a NOVX protein in humans and for treating a pathology associated with the human disease. NOVX nucleic acids, proteins and CC antibodies can be used in the treatment and diagnosis of cardiomyopathy, CC atherosclerosis, hypertension, congenital heart defects, aortic stenosis, CC valve disease, tuberous sclerosis, scleroderma, obesity, transplantation, CC disorders, neoplasm, lymphoma, uterus cancer, diabetes, metabolic disease, alds, brochial asthma, Crohn's disease, multiple sclerosis, C haematopoietic disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, anorexia, cancer-associated cachexia, cancer, C Alzheimer's disease, Parkinson's disease, immune disorders, C ACC62346 to ACC62465 represent PCR primers and metabolic syndrome X. Sequences, which are used in examples from the present invention. With the human NOV35b protein in the exemplification of the present invention. Sequence 520 AA;

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The invention discloses a composition comprising two or more isolated rat or human polynucleotides or a polynucleotide which represents a fragment, derivative or allelic variation of the nucleic acid sequence. Also claimed are a vector comprising the novel polynucleotide, a host cell comprising the vector, a method for identifying a nucleotide sequence which is differentially regulated in an animal subjected to pain and a kit to perform the method, an array, a method for identifying an agent that increases or decreases the expression of the polynucleotide sequence that is differentially expressed in neuronal tissue of a first animal subjected to pain, a method for identifying an agent the expression of a polynucleotide sequence which is differentially expressed in an animal subjected to pain, a method for identifying a compound which regulates the activity of one or more of the polynucleotides, a method for producing a pharmaceutical composition, a method for identifying a compound useful in treating specification, a method for identifying a compound useful in treating pain and a pharmaceutical composition comprising the one or more goain (e.g. spinal segmental nerve injury (Chung), chronic constriction injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene therapy). The sequence presented is a human protein (shown in Table 2 of
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The invention discloses a composition comprising two or more isolated rat or human polynucleotides or a polynucleotide which represents a fragment, clearly at the or allelic variation of the nucleic acid sequence. Also claimed are a vector comprising the novel polynucleotide, a host cell comprising the vector, a method for identifying a nucleotide sequence which is differentially regulated in an animal subjected to pain and a compound that increases or decreases the expression of the polynucleotide sequence that increases or decreases the expression of the polynucleotide sequence that is differentially expressed in neuronal tissue of a first animal subjected to pain, a method for identifying a compound which regulates the expression of a polynucleotide sequence which is differentially expressed in an animal subjected to pain, a method for identifying a compound that regulates the activity of one or more of the polynucleotides, a method for producing a pharmaceutical composition, a method for identifying a compound useful in treating pain and a pharmaceutical composition comprising the one or more of pain, and a pharmaceutical composition of the polypeptides given in the specification, a method for identifying a compound useful in treating pain and a pharmaceutical composition comprising the one or more of polypeptides or their antibodies. The polynucleotide or the compound that composition is useful for preparing a medicament for treating pain and a pharmaceutical nerve injury (CNung), chronic constriction or injury (CCI) and spared nerve injury (SNI) in an animal (e.g. gene of the specification) which is differentially expressed during pain. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic form directly from WIPO at the printed specification at the pain and the printed contributed the printed specification at the printed specif
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High-efficiency full-length cDNA cloning
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Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
S Carninci, P., Shibata, Y., Muramatsu, M. and Hayashizaki, Y.

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genome Res. 10 (10), 1617-1630 (2000)

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Group Phase I & II Team.

Analysis of the mouse transcriptome based on functional anno of 60,770 full-length cDNAs

Nature 420, 563-573 (2002)

6 (bases 1 to 3286)

Adachi,J., Aizawa,K., Akimura,T., Arakawa,T., Bono,H., Carni Fukuda,S., Furuno,M., Hanagaki,T., Hara,A., Hashizume,W., Hayashida,K., Hayatsu,N., Hiramoto,K., Hiraoka,T., Hirozane, Hori,F., Imotani,K., Ishii,Y., Itoh,M., Kagawa,I., Kasukawa, Katoh,H., Kawai,J., Kojima,Y., Kondo,S., Konno,H., Kouda,M., Koya,S., Kurihara,C., Matsuyama,T., Miyazaki,A., Murata,M.,
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AK044835.1 GI:26336856
HTC; CAP trapper.
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FANTOM Consortium.
Functional annotation of a full-length mouse
Nature 409, 685-690 (2001)
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The FANTOM Consortium and the RIKEN Genome Ex
Group Phase I & II Team.
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High-efficiency full-length cDNA cloning
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Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Kanagawa 230-0045, Japan (B-mail:genome-res@gsc.riken.go.jp, Fax:81-45-503-9216)

CDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Exploration Research Group in Riken Division of Experimental Animal Research in Riken contributed to Fissues were provided by Dr. Tomohiro Kono (Department of Animal Science, Tokyo University of Agriculture, 1737 Hunako Atsugi City, acknowledge.
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Please visit our web site for further URL:http://genome.gsc.riken.go.jp/
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Carninci, P. and Hayashizaki, Y.

High-efficiency full-length cDNA competh. Enzymol. 303, 19-44 (1999)

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Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

of 60,770 full-length cDNAs

Nature 420, 563-573 (2002)

E 6 (bases 1 to 3466)

E Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W., Hayashida, K., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kasukawa, T., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Murata, M., Nakamura, M., Nishi, K., Nomura, K., Numazaki, A., Murata, M., Ohazaki, Y., Saitoh, H., Sakai, C., Sakai, K., Sakazume, N., Sano, H., Sasaki, D., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Tagami, M., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Takeda, Y., Tanaka, T., Tomaru, A., Toya, T., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     prepare full-length cDNA libraries for rapid discovery of new genes Genome Res. 10 (10), 1617-1630 (2000)

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E 3

S Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P., Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M., Sumi, N., Ishii, Y., Nakamura, S., Hazama, M., Nishine, T., Harada, A., Yamamoto, R., Matsumoto, H., Sakaguchi, S., Ikegami, T., Kashiwagi, K., Fujiwake, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y. RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer

L Genome Res. 10 (11), 1757-1771 (2000)

E 1076861
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                                                                                                                                                                                                                                                                                                                                        Direct Submission
Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GS RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yok Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp, URL:http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222, Fax:81-45-503-9216)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           The RIKEN Genome Exploration Research Group F FANTOM Consortium.
Functional annotation of a full-length mouse Nature 409, 685-690 (2001)
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Please visit our web site for further details.
                                                                                                                                                                                             The FANTOM Consortium and the RIKEN Genome Exploration Research
     /tissue
/clone_ī
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                                                                                 /mol_type="mRNA"
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Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Lou Staudt
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information car
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM12877 row: h column: 07
High quality sequence stop: 595.
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National Institutes of Health,
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Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: James Cleaver, M.D.
cDNA Library Preparation: Life Technologies, Ir
cDNA Library Arrayed by: The I.M.A.G.E. Consort
Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution info
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM10700 row: c column: 21
High quality sequence stop: 815.
Location/Qualifiers
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/clone_lib="NCI_CGAP_Skn3"
/note="Organ: skin; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: SalI; Cloned unidirectionally. Primer: Oligo dT
Average insert size 1.5kb. Library constructed by Life
Technologies. Note: this is a NCI_CGAP_Library."
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Email: cgapbs-r@mail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: Rubin Laboratory

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Agencourt Bioscience Corporation

Clone distribution: MGC clone distribution information car

found through the I.M.A.G.E. Consortium/LLNL at:

http://image.llnl.gov

plate: LLCM2139 row: p column: 16

High quality sequence start: 3

High quality sequence stop: 693.

Location/Qualifiers

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Mammalia; Eutheria; Primates; Cat
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NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Ma
Unpublished (1999)
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/mol_type="mRNA"
/mol_type="mRNA"
/db_xref="taxon:9606"
/db_xref="taxon:9606"
/clone="IMAGE:5952255"
/clone="IMAGE:5952255"
/tissue_type="ductal carcinoma, cell line"
/tissue_type="ductal carcinoma, cell line"
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/clone_lib="NIH_MGC_110"
/clone_lib="NIH_MGC_110"
/note="Organ: pancreas; Vector: pOTB7; Site_1: XhoI;
Site_2: EcoRI; cDNA made by oligo-dT priming.
Site_3: About a site of the state of the state of the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies).
Note: this is a NIH_MGC Library."
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Pred. No. 2.3e-112;
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Tissue Procurement: James Cleaver, M.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information car
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM10629 row: f column: 23
High quality sequence stop: 794.
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NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Ma
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: James Cleaver
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National Institutes of Health, Mammalian
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
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Tissue Procurement: James Cl
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/db_xref="taxon:9606"
/clone="IMAGE:4778093"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NCI_CGAP_Skn3"
/note="Organ: skin; Vector: pCMV-SPORT6; Site_1: NotI; Site_2: SalI; Cloned unidirectionally. Primer: Oligo Average insert size 1.5kb. Library constructed by Lif Technologies. Note: this is a NCI_CGAP_Library."
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1 (bases 1 to 894)

NIH-MGC http://mgc.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC Unpublished (1999)

Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Ling Hong/Rubin Laboratory
cDNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can befound through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLCM1614 row: h column: 10
High quality sequence stop: 799.
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/clone_lib="NHI_MGC_47"
/clone_lib="NHI_MGC_47"
/cloned_ib="NHI_MGC_47"
/cloned_into_EcoRI/XhoI sites using the following 5'
cloned_into_EcoRI/XhoI sites using the following 5'
adaptor: GGCACGAG(G). Size-selected >500bp for average
insert_size_1.8kb. Library_constructed_by_Ling_Hong_in
the_laboratory_of_Gerald_M. Rubin_(University_of_
California, Berkeley)_using_ZAP-cDNA_synthesis_kit
(Stratagene)_and_Superscript_II_RT_(Life_Technologies).
Note: this is a NIH_MGC_Library."
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Ota, T., Nishikawa, T., Su
Yamamoto, J., Wakamatsu, A
Isogai, T.
HRI human cDNA project
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Homo sapiens
Eukaryota; Metazoa; C
Mammalia; Eutheria; P
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AU127534
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1532-3 Yana, Kisarazu, Chiba 292-0812, Japan
Tel: 81-438-52-3975
Fax: 81-438-52-3986
Email: genomics@hri.co.jp
HRI human cDNA project; 5'- & 3'-end one pass sequencing:
Research Institute; cDNA library construction: Department
Virology, Institute of Medical Science, University of Toky
Helix Research Institute.
Location/Qualifiers
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Genomics Laboratory
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ilarity 97.0%;
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Eutheria;
1 to 703)
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/db_xref="taxon:9606"
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                                                                                                                                                                                                                                                                                                /cell_type="teratocarcinoma"
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/clone_lib="NT2RP2"
/note="Vector: pME18SFL3; mRNA from cells after 2-weeks retinoic acid (F
                                                                                                                                                                                                                                                                                                                                                                                                organism="Homo"
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u,A., Nakamura,Y.,
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ıra,Y., Nagai,T., Su
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DX BX441322 Homo sapiens FETAL BRÄIN Homo sapiens cDNA clone CSODFO16YJ04 5-PRIME, mRNA sequence.

RX441322

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BX441322.1 GI:30789948

EST.

Homo sapiens (human)

SM Homo sapiens (human)

SM Homo sapiens (human)

E Lukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

E 1 (bases 1 to 1083)

S Li,W.B., Gruber, C., Jessee, J. and Polayes, D. Full-length cDNA libraries and normalization Unpublished (2001)

Contact: Genoscope

Genoscope - Centre National de Sequencage

BP 191 91006 EVRY cedex - France

Email: seqref@genoscope.cns.fr, Web: www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of Invitrogen. This sequence belongs to sequence cluster 3098.r For more information about this cluster, see

http://www.genoscope.cns.fr

http://fulllength.invitrogen.com/ InvitroGen Corporation 1600

Faraday Avenue Genoscope sequence ID: CSODFO16DE02QP1.

Location/Qualifiers

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/db_xref="taxon:9606"

/clone="CSODF016YJ04"

/tissue_type="FETAL BRAIN"

/dev_stage="fetal"

/clone_lib="Homo sapiens FETAL BRAIN"

/note="Organ: brain; Vector: pCMVSPORT_6; lst strand cDNA

was primed with a NotI-oligo(dT) primer. Five prime end
enriched, double-strand cDNA was digested with Not I and
cloned into the Not I and EcoRV sites of the pCMVSPORT 6

vector. Library was not normalized."
                                                                                                            /organism="Homo
/mol_type="mRNA"
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                                                                                                      GGATGTTGCTGAATGACTTCCTTAATGACCAGAATAGAGATAAAAATAGTATCTATAGTG
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Mammalia; Eutheria; Primates; Cat
1 (bases 1 to 795)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Ma
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
 496
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High quality sequence stop: 758.
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GAAAACATGAAGCTCAGAGAGGACACCCATCATTTGGTCAACTATATTCCCCAGGGTTCA
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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4761657"
/tissue_type="neuroblastoma, cell line"
/lab_host="DH10B (phage-resistant)"
/clone_lib="NIH_MGC_47"
/clone_lib="NIH_MGC_47"
/note="Organ: brain; Vector: pOTB7; Site_1: XhoI; Site_2:
cloned_into_EcoRI/XhoI sites_using_the_following_5'
adaptor: GGCACGAG(G). Size-selected >500bp for average
insert_size_1.8kb. Library_constructed_by_Ling_Hong_in
the_laboratory_of_Gerald_M. Rubin_(University_of_California, Berkeley) using_ZAP-cDNA_synthesis_kit
(Stratagene)_and_Superscript_II_RT_(Life_Technologies).
Note: this is_a_NIH_MGC_Library."
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: James Cleaver, M.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LL
Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM10694 row: l column: 23
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Eukaryota; Metazoa; Chordata; Cra
Mammalia; Eutheria; Primates; Cat
1 (bases 1 to 733)
NIH-MGC http://mgc.nci.nih.gov/
National Institutes of Health, Ma
Unpublished (1999)
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602658594F2 NCI_CGAP_9
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BG698557.1 GI:1396599
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IMAGE:4801702 5',
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Rattus norvegicus (Norway ra
Rattus norvegicus
Eukaryota; Metazoa; Chordata
Mammalia; Eutheria; Rodentia
Rattus.
1 (bases 1 to 789)
Bonaldo, M.F., Lennon, G. and
Normalization and subtractic
discovery
Genome Res. 6 (9), 791-806
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Sciurognathi; Muridae; Murinae;
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Best Local Similarity 88.3%;
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Coordinated Laboratory f
University of Iowa
375 Newton Road , 4156
Tel: 319 335 8250
Fax: 319 335 9565
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8889548
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Tissue Procurement: Dr. James Lin, Universtiy of Iowa
cDNA Library preparation: Dr. M. Bento Soares, University of
cDNA Library Arrayed by: Dr. M. Bento Soares, University of
DNA Sequencing by: Dr. M. Bento Soares, University of
Clone Distribution: Researchers may obtain clones from Rese
Genetics (www.resgen.com).
Seq primer: M13 REVERSE.
Location/Qualifiers
                                                                                                                                                                                                                                                                                             AAAATAGTATCTATAGTGGCCTGGAAGCCTTTGGGGATGTTAAATTAGAAGACACCTACT
                                       CAAAGGCATCTTTACTTGTATCAAATCAAAATGGAAATATGTACACATCTTCAGTATATG
                                                                                                    TGCAGAAATCTCTAGCTAGGATGTTCCTGAATGACTTTCTTAACGATCAAAACAGAGACA
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          GTTCCCTTGCATCTGTTCTAGCACAGTACTCACCTCAGCAATTAGCAGGGAAGAGAATTG
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/strain="Sprague-Dawley"
/db_xref="taxon:10116"
/clone="UI-R-FJO-cpz-j-19-0-UI"
/tissue_type="embryo"
/dev_stage="embryo"
/dev_stage="embryo"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/clone lib="UI-R-FJO"
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http://fulllength.invitrogen.com/ Faraday Avenue Genoscope sequence Location/Qualifiers 11201 /organism="Homo sapiens" /mol_type="mRNA" /clone="CSODF022YE18" /tissue_type="FETAL BRAIN /dev_stage="fetal" /clone_lib="Homo sapiens /note="Organ: brain; Vectowas primed with a NotI-olenriched, double-strand cocloned into the Not I and	1 91006 EVRY cedex - France segref@genoscope.cns.fr, Web : wry was constructed by Life Technol rogen. This sequence belongs to se information about this cluster, se //www.genoscope.cns.fr/ in/cluster.cgi?seq=CSODF022BC09QP1 in/cluster.cgi?seq=CSODF022BC09QP1	bases 1 to 1201) .B., Gruber,C., Jessee,J. and Polayes,Dlength cDNA libraries and normalization blished (2001) act: Genoscope scope - Centre National de Sequencage	EST. Homo sapiens (human) Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata;	BX419944 Homo sapiens FETAL BRAIN Homo sapiens cDNA clone CSODF022YE18 5-PRIME, mRNA sequence. BX419944 GI:30650862	490 AGCATATTC 1498 781 AGCATATTC 789	1430 CTCCAAATGATGACACTTTGGATGAAGGAGTAGGACTTGTGCATTCAAACATAGCAACTG 1489 	.370 GAACGTGGTACTTAGTTAGGGTGGATGAAAAGCACAGAAGAACTTACGCTCGGCGTCCCA 1429 	.310 AGGACACCCATCATTTGGTCAACTATATTCCCCCAGGGTTCAATAGATTCACTCTTTGAAG 1369	.250 GGCTTGATTCAAGAACTGGTGTGGCACCAGATGTCTTCGCTGAAAACATGAAGCTCAGAG 1309 	190 ATGCTACACCGGGGTCTGCTCTTGATAAAATAACAGCAAGTTTATGTGATCTTAAATCAA 1249 	130 GAGIGITITCTIAIGGITCIGGITIGGCIGCCACICTGIACTCICITAAAGICACACAAG 1189	
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999 TWAGARGCACCTCT 1012

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Search completed: June 24, 2004, 11:14:15 Job time : 4695 secs

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-MODEL=frame+_p2n.model -DEV=xlp
-Q=/cgn2_1/USPTO_spool_p/US10622516/runat_23062004_162633_648/app_query.fas
-DB=GenEmbl -QFMT=fastap -SUFFIX=rge -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pto -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
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-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAP(-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7
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1 (bases 1 to 2068)

S trausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G. Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K. Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J., Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J., WcKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., WcKernan, K.J., Hulyk, S.W., Villalon, D.K., Marny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A., Fahey, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S., Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G., Blakesley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.N. Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E., Schnerth, A., Schein, J.E., Jones, S.J. and Marra, M.A. Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences before than 15,000 full-length human and mouse cDNA sequences and mouse cDNA sequences and more than 15,000 full-length public of the Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590.
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NIH-MGC Project URL: http://mgOn Aug 20, 2003 this sequence Contact: MGC help desk Email: cgapbs-r@mail.nih.gov Tissue Procurement: DCTD/DTP cDNA Library Preparation: Rubi
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Shenmen,C.M., Schuler,G.D.,
Schaefer,C.F., Bhat,N.K.,
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Web site: http://www.nisc.nih.gov/
Contact: nisc_mgc@nhgri.nih.gov
Akhter,N., Ayele,K., Beckstrom-Sternberg,S.M., Benjamin,B.,
Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C., Brooks,S.,
Dietrich,N.L., Granite,S., Guan,X., Gupta,J., Haghighi,P.,
Hansen,N., Ho,S.-L., Karlins,E., Kwong,P., Laric,P., Legaspi,R.,
Maduro,Q.L., Masiello,C., Maskeri,B., Mastrian,S.D.,McCloskey,J.C.,
McDowell,J., Pearson,R., Stantripop,S., Thomas,P.J., Touchman,J.W.,
Tsurgeon,C., Vogt,J.L., Walker,M.A., Wetherby,K.D., Wiggins,L.,
Young,A., Zhang,L.-H. and Green,E.D.
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This clone was selected for full length sequencing because passed the following selection criteria: matched mRNA gi: 'Docation/Qualifiers 1.2068
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/note="synonym: HMGCS"
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Hydroxymethylglutaryl-coenzyme
/db_xref="CDD:pfam01154"
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/gene="HMGCS1"
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Qy 179 SerileGlnCysTyrLeuSerAlaLeuAspArgCysTyrSerValTyrCysLysIle 198	663 ATTGGGCCAAATGCTCCTTTAATTTTTGAACGAGGCTTCGTGGGACACATATGCAACAT 72 159 AlaTyrAspPheTyrLysProAspMetLeuSerGluTyrProIleValAspGlyLysLeu 17	GTGGAGTTGGAGCAGTAGCTCTGCT	Qy 141 AsnTrpIleGluSerSerSerTrpAsp	Qy 121 GluGlyIleAspThrThrAsnAlaCysTyrGlyGlyThrAlaAlaValPheAsnAlaVal 140 	Qy 101 SerLysSerValLysThrAsnLeuMetGlnLeuPheGluGluSerGlyAsnThrAspIle 120 	nLeuSerTyrAspCysIleGlyArgLeuGluVa. { CCTTTCCTATGATTGCATTGGGCGGCTGGAAGTT	Oy 61 AspArgGluAspIleAsnSerLeuCysMetThrValValGlnAsnLeuMetGluArgAsn 80 	Qy 41 GlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMetGlyPheCysThr 60 	AspGlnAlaGluLeuc ATCAAGCAGAGTTGC	1 MetProGlySerLeuProLeuAsnAl	uery Match: 97.13% Indels: B: 97.13% Gaps: S-10-622-516-2 (1-478) x HSCOAS (1-1685)	Baca	RIGIN EPEAAVISNGVW"	NSIYSGLKAFG YGSLASVLAQY LKSRLDSRTGV	GKYTIGLGQAKMGFCTDREDINSLCMTVVQNLMERNNLSYDCIGRLEVGTETIIDKSK SVKTNLMQLFEESGNTDIEGIDTTNACYGGTAAVFNAVNWIESSSWDGRYALVVAGDI AVYATGNARPTGGVGAVALLIGPNAPLIFERGLRGTHMQHAYDFYKPDMLSEYPIVDG KLSIOCYLSALDRCYSVYCKKIHAOWOKEANDNDFTLNDFGFMIFHSPYCKLVOKSLA
, Takahashi-Fujii,A., Tanase,T. Musashino,K., Yuuki,H., Hara,H, Sato,H., Ota,T., Wakamatsu,A. Kawai-Hio,Y., Saito,K., Nishik,H., Matsuo,K., Nakamura,Y., Se Wagatsuma,M., Murakawa,K., Kan B., Suzuki,Y., Sugano,S., Nagah i,T.	RSION AK0954 YWORDS oligo URCE Homo s ORGANISM Homo s Eukary Mammal	AK09549 DN Homo sa to HYDR		9 ValGlyLeuValHisSerAsnIleAlaThrGluHisIleProSerProAlaLysLys	419 LysHisArgArgThrTyrAlaArgArgProThrProAsnAspAspThrLeuAspGluGly 43	399 ProGlnGlySerIleAspSerLeuPheGluGlyThrTrpTyrLeuValArgValAspGlu 41	Qy 379 AspValPheAlaGluAsnMetLysLeuArgGluAspThrHisHisLeuValAsnTyrIle 398	359 IleThrAlaSerLeuCysAspLeuLysSerArgLeuAspSerArgThrGlyValAlaPro 37	9 AlaThrLeuTyrSerLeuLysValThrGlnAspAlaThrProGlySerAlaI	Qy 319 SerProGlnGlnLeuAlaGlyLysArgIleGlyValPheSerTyrGlySerGlyLeuAla 338	Qy 299 AsnGlyAsnMetTyrThrSerSerValTyrGlySerLeuAlaSerValLeuAlaGlnTyr 318	79 LysAlaSerSerGluLeuPheSerGlnLysThrLysAlaSerLeuLeuVals 	Qy 259 PheGlyAspValLysLeuGluAspThrTyrPheAspArgAspValGluLysAlaPheMet 278	Qy 239 AsnAspPheLeuAsnAspGlnAsnArgAspLysAsnSerIleTyrSerGlyLeuGluAla 258	Qy 219 MetIlePheHisSerProTyrCysLysLeuValGlnLysSerLeuAlaArgMetLeuLeu 238

41 GlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMetGlyPheCysThr 6	Length: 3195 Score: 2403.50 Matches: 467 Percent Similarity: 89.81% Conservative: 0 Best Local Similarity: 89.81% Mismatches: 0 Query Match: 95.72% Indels: 53 DB: US-10-622-516-2 (1-478) x AK095492 (1-3195) Qy	/clone_lib="FCBBF1" //dev_stage="fetal" /note="cloning vector: pME18SFL3" 681597 /note="unnamed protein product" /codon_start=1 /protein_id="BAC04559.1" /db_xref="GI:21754758" /translation="MPGSLPLNAEACWPKDVGIVAL GKYTIGLGQAKMGFCTDREDINSLCMTVVQNLMERN SVKTNLMQLFEESGNTDIEGIDTTNAVNWIESSSWD GGVGAVALLIGPNAPLIFERGLRGTHMQHAYDFYKP DRCYSVYCKKIHAQWQKEGNDKDFTLNDFGFMIFHS NRDKNSIYSGLEAFGDVKLEDTYFDRDVEKAFMKAS TSSVYGSLASVLAQYSPQQLAGKRIGVFSYGSGLAA SLCDLKSRLDSRTGVAPDVFAENMKLREDTHHLVNY HRRTYARRPTPNDDTLDEGVGLVHSNIATEHIPSPA H" RIGIN	search Association for Biotechnology (RAB); cDNA library nstruction: Helix Research Institute (HRI) (supported by Japan y Technology Center etc.); 5'- & 3'-end one pass sequencing: RAI, and Biotechnology Center, National Institute of Technology a aluation; clone selection for full insert sequencing: HRI and B; annotation: HRI and RAB. Location/Qualifiers 13195 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="FCBBF1000053" /tissue type="brain"	EDO human cDNA sequencing project npublished (bases 1 to 3195) sogai,T. and Yamamoto,J. irect Submission ubmitted (04-JUL-2002) Takao Isogai, FLJ Project(azusa-Kamatari, Kisarazu, Chiba 292-0812, Japan azusa-Kamatari, Kisarazu, Chiba 292-0812, Japan E-mail:genomics@hri.co.jp, Tel:81-438-52-3975, FaEDO human cDNA sequencing project supported by Miconomy, Trade and Industry of Japan; cDNA full in
Qy 359 IleThrAlaSerLeuCysAspLeuLysSerArgLeuAspSerArgThrGlyValAlaPro 378	Oy 279 LysalaSerSerGluLeuPheSerGlnLysThrLysalaSerLeuLeuValSerAsnGln 298	179 SerileGlnCysTyrLeuSerAlaLeuAspArgCysTyrSerValTyrCysLysIle	ACTGGATTGAGTCCAGCTCTTG CTGTATATGCCACAGGAAATGC TTGGGCCAAATGCTCTTAAT ATYXASPPheTYXLYSPYOAS ATYXASPPheTYXLYSPYOAS ATYXASPPHETYXLYSPYOAS ATTGATGATTTACAAGCCTGA	Qy 101 SerLysSerValLysThrAsnLeuMetGlnLeuPheGluGluSerGlyAsnThrAspIle 120

Qy 121 GluGlyIleAspThrThrAsnAlaCysTyrGlyGlyThrAlaAlaValPheAsnAlaV	Qy 101 SerLysSerValLysThrAsnLeuMetGlnLeuPheGluGluSerGlyAsnThrA	Qy 81 AsnLeuSerTyrAspCysIleGlyArgLeuGluValGlyThrGluThrIleIleAs: :::	Qy 61 AspArgGluAspIleAsnSerLeuCysMetThrValValGlnAsnLeuMetGluA	GlyGlnAlaL; : GGCCAGGCCA	aLeuGluIleTyrPheProSerGlnTyrValAspGlnAlaGluLeuG	1 MetProGlySerLeuProLeuAsnAlaG 	ry Match: 94.07% Indels: 4 10-622-516-2 (1-478) x AX700129 (1-3275)	ed. No.: 4.77e-192 Length ore: 2362.00 Matche rcent Similarity: 89.62% Conser st Local Similarity: 87.50%	IGIN icoment Scores.	/organism="Rattus norvegicus" /mol_type="unassigned DNA" /db_xref="taxon:10116" /note="Cytosolic 3-hydroxy 3-methylclutary]	EP 1284298-A 15 19-FEB-2003; LAMBERT COMPANY (US) Location/Qualifiers 1 3275	ERENCE 1 UTHORS Brooksbank, R.A., Dixon, A.K., Lee, K ITLE Identification and use of molecule	ORGANISM Rattus norvegicus Eukaryota; Metazoa; Chordata Mammalia; Eutheria; Rodentia	RSION AX700129.1 G YWORDS .	AX700129 DN Sequence 15 from Patent EP1284298. V AX700129	SULT 7 700129	Qy 459 ProArgLeuProAlaThrAlaAlaGluProGluAlaAlaValIleSerAsnGlyG	Qy 439 ValGlyLeuValHisSerAsnIleAlaThrGluHisIleProSerProAlaLysLy	
laVal 140 Qy 43	Aspīle 120 GATATA 451	pLys 100 CAAA 391	rgAsn 80 GAAAT 331	yeThr 60 CACG 271	Asp 40 Db 12 GAT 211	leval 20 CGTT 151	Qy 29 Db 111	Qy 27 Db 105		Ортууль в СОртууль в 1 др 93	Оу 21 рь 87	Db 81	omi;	Ov 17	03-APR-2003 Qy 15	;	GluHis 478 GAACAT 1594	sVal 458 AGTA 1534	 AAGGA 1474
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Qy 149 149	Qy 141 AsnTrpIleGluSerSerSerTrpAsp 149	Qy 121 GluGlyIleAspThrThrAsnAlaCysTyrGlyGlyThrAlaAlaValPheAsnAlaVal 140	Qy 101 SerLysSerValLysThrAsnLeuMetGlnLeuPheGluGluSerGlyAsnThrAspIle 120	Qy 81 AsnLeuSerTyrAspCysIleGlyArgLeuGluValGlyThrGluThrIleIleAspLys 100 :::	Qy 61 AspArgGluAspIleAsnSerLeuCysMetThrValValGlnAsnLeuMetGluArgAsn 80	QY 41 GlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMetGlyPheCysThr 60	Qy 21 AlaLeuGluIleTyrPheProSerGlnTyrValAspGlnAlaGluLeuGluLysTyrAsp 40	QY 1 MetProGlySerLeuProLeuAsnAlaGluAlaCysTrpProLysAspValGlyIleVal 20	· Gaps:	0500 0000 0000 0000	ignment Scores: 4.77e-192 Length: 3	type="unassigned DNA" ref="taxon:10116"	cion/Qualifiers 275 nism="Rattus norvegi	TITLE Methods for the toxicity pr JOURNAL Patent: EP 1344834-A 37 17- F. HOFFMANN-LA ROCHE AG (CH	Eukaryota; Metazoa; Chordata; Cra Mammalia; Eutheria; Rodentia; Sci Rattus.	YWORDS URCE Rattus norvegicus (Norway rat) ORGANISM Rattus norvegicus	CUS AX827303 FINITION Sequence 37 from Paten CESSION AX827303	SULT 8	PIOATGLEUPTOALAThTALAALAGIUPTOGIUAlaAlaValIleSerAsnGlyGluHis 478	ASO Drobroklambralasisci
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9 RNCHMCA 3275 bp mRNA linear ROD 18-JUN-2003	459 ProArgLeuProAlaThrAlaAlaGluProGluAlaAlaValIleSerAsnGlyGluHis 478	439 ValGlyLeuValHisSerAsnIleAlaThrGluHisIleProSerProAlaLysLysVal 458		399 ProGlnGlySerIleAspSerLeuPheGluGlyThrTrpTyrLeuValArgValAspGlu 418	379 AspValPheAlaGluAsnMetLysLeuArgGluAspThrHisHisLeuValAsnTyrIle 398	359 IleThrAlaSerLeuCysAspLeuLysSerArgLeuAspSerArgThrGlyValAlaPro 378	339 AlaThrLeuTyrSerLeuLysValThrGlnAspAlaThrProGlySerAlaLeuAspLys 358	319 SerProGlnGlnLeuAlaGlyLysArgIleGlyValPheSerTyrGlySerGlyLeuAla 338		ω υ	992 TTTGGGGATGTGAAATTAGAAGATACTTACTTCGACAGAGATGTGGAAAAGGCATTTATG 1051	259 PheGlyAspValLysLeuGluAspThrTyrPheAspArgAspValGluLysAlaPheMet 278	239 AsnAspPheLeuAsnAspGlnAsnArgAspLysAsnSerIleTyrSerGlyLeuGluAla 258	219 MetilePheHisSerProTyrCysLysLeuValGlnLysSerLeuAlaArgMetLeuLeu 238 	199 HisAlaGlnTrpGlnLysGluGlyAsnAspLysAspPheThrLeuAsnAspPheGlyPhe 218 	179 SerIleGlnCysTyrLeuSerAlaLeuAspArgCysTyrSerValTyrCysLysLysIle 198	159 AlaTyrAspPheTyrLysProAspMetLeuSerGluTyrProIleValAspGlyLysLeu 178 	D	50GlyLeuArgGlyThrHisMetGlnHis 15	572 GCTATATATGCCTCAGGAAACGCCAGGCCTACAGGTGGAGTTGGAGCTGTGGCTCTGCTA 631

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of Biochemistry, School of
barcelona, Spain
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Cytosolic 3-hydroxy
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Nucleotide sequence of a rat liver cDNA encoding
3-hydroxy-3-methylglutaryl coenzyme A synthase
Nucleic Acids Res. 18 (12), 3642 (1990)
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RMFLNDFLNDQNRDKNSIYSGLEAFGDVKLEDTYFDRDVEKAFMKASAELFNQKTKAS
LLVSNQNGNMYTSSVYGSLASVLAQYSPQQLAGKRIGVFSYGSGLAATLYSLKVTQDA
TPGSALDKITASLCDLKSRLDSRTCVAPDVFAENMKLREDTHHLANYIPQCSIDSLFE
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Hamilia Lia; Bulletia; Noterita; Scrutygratia; Fullicas; Fullicas; Numbers; R.L., Peingold, E.A., Grouse, L.H., Derge, J.G.,
Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
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E. 2 (bases 1 to 2703)
Strausberg, R.
Direct Submission
L. Submitted (05-FEB-2002) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
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Mammalia; Eutheria;
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Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Jeffrey Green M.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc_mgc@nhgri.nih.gov

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McDowell,J., Pearson,R., Stantripop,S., Thomas,P.J., Touchman,J.W.,
Tsurgeon,C., Vogt,J.L., Walker,M.A., Wetherby,K.D., Wiggins,L.,
Young,A., Zhang,L.-H. and Green,E.D.
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This clone was selected for full length sequencing because passed the following selection criteria: matched mRNA gi: incomplete incomplete continuous   (1-
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  /protein_id="AAH23851.1"
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KLSIQCYLSALDRCYSVYRKKIRAQWQKEGKDKDFTLNDFGFMIFHSPYCKLVQKSLA
RMFLNDFLNDQNRDKNSIYSGLEAFGDVKLEDTYFDRDVEKAFMKASSELFNQKTKAS
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GTWYLVRVDEKHRRTYARRPFTNDHSLDEGMGLVHSNTATEHIPSPAKKVPRLPATSA
ESESAVISNGEH"
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/lab_host="DH10B"
/note="Vector: pCMV-SPORT6"
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/note="HMG_CoA_synt; Region:
Hydroxymethylglutaryl-coenzyme
/db_xref="CDD:pfam01154"
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         Eukaryota;
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                                                                                                    musculus
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                                               3202 bp mRNA linear 3-hydroxy-3-methylglutaryl-Coenzyme A MGC:36662 IMAGE:5366786), complete cds
                                      GI:21706865
                           (house
          Chordata;
Rodentia;
                           mouse)
          Craniata; Vertebrata;
Sciurognathi; Muridae;
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synthase 1,
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; Murinae; Mus
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Oy 279 LYSALaSerSerGluLeuPheSerGlnLysThrLysAlaSerLeuLeuValSerAsnGln 298	AAGGATAAAGATTTTACCCTGAATGATTTTGGCTTC LYSLeuValGlnLysSerLeuAlaArgMetLeuLeu	565 ATTGGGCCAAACGCTCCTCTAATTTTTGACCGAGGGCTCCGTGGGACACACAC	heGluGluSerGlyAsnThrAspile 1	Qy 21 AlaLeuGluIleTyrPheProSerGlnTyrValAspGlnAlaGluLeuGluLysTyrAsp 40

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S Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.I.
Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.
Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F.,
Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L.,
Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S.,
Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,
Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J.,
McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S.,
Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,
Villalon, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A.,
Fahey, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S.,
Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y.,
Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D.,
Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M.
Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E.,
Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.
Generation and initial analysis of more than 15,000 full-length
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human and mouse cDNA sequence. Natl. Acad. Sci. U. 22388257
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2 (bases 1 to 3260)
Strausberg, R.
Direct Submission
Submitted (06-MAY-2002) N
Gene Collection (MGC), Calinstitute, 31 Center DrivusA
NIH-MGC Project URL: http.
Contact: MGC help desk
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Tissue Procurement: Gilbert Smith, Ph.D.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LL)
DNA Sequencing by: Baylor College of Medicine Human Ge
Sequencing Center
Center code: BCM-HGSC
Web site: http://www.hgsc.bcm.tmc.edu/cdna/
Contact: amg@bcm.tmc.edu
Gunaratne, P.H., Garcia, A.M., Lu, X., Hulyk, S.W., Lo
Kowis, C.R., Sneed, A.J., Martin, R.G., Muzny, D.M., N
A.N., Gibbs, R.A.
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This clone was selected for full length sequencing because passed the following selection criteria: matched mRNA gi: incomplete in the contion of the continuous continuous continuous distribution of the continuous conti
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ESESAVISNGEH"
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EMERTYOTA; Metazoa; Chordata; Craniata, Vertebrata; Euteleostomi;

Elkaryota; Metazoa; Chordata; Craniata, Vertebrata; Euteleostomi;

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1 (bases 1 to 3278)

21 (bases 1 to 3278)

22 1 (bases 1 to 3278)

23 Strausberg, R. L., Feingold, E.A., Grouse, L.H., Derge, J.G.,

Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,

Holkins, R.F., Jordan, H., Moore, T., Max, S.I., Mang, J., Hsieh, F.,

Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,

Stapleton, M., Soares, M. B., Bonaldo, M.F., Casvant, T.L.,

Scheetz, T.E., Brownstein, M. J., Uddin, T. B., Toshiyki, S.,

Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J.,

Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.L.,

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Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W.,

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Fahey, J., Helton, E., Ketteman, M., Madan, A., Rodrigues, S.,

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Butterfield, Y.S., Kzywinski, M.I., Skaiska, U., Smailus, D.E.,

Schmerch, A., Schein, J.E., Jones, S.J., Lu, X., Grimusod, J., Wers, R.M.,

Butterfield, Y.S., Kzzywinski, M.I., Skaiska, U., Smailus, D.E.,

Schmerch, A., Schein, J.E., Jones, S.J., and Mazra, M.A.

Gene Collection and initial analysis of more than 15,000 full-length human and mouse cOMA sequences

L. Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)

238257

25 (bases I. to 3278)

Strausberg, R.

Direct Submission

ALL Gene Collection (MGC), Cancer Genomics office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA

Contact: MGC help desk

Email: cagabs-remail.nih.gov

Tissue Procurement: The Cepko Laboratory

CDNA Library Preparation: Life Technologies, Inc.
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This clone was selected for full length sequencing because passed the following selection criteria: matched mRNA gi: []
GlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMetGlyPheCysThr
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                                      AlaLeuGluIleTyrPhePro
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Alignment Scores: Pred. No.: 2089.00 Matches: Percent Similarity: Best Local Similarity: 83.33% Query Match: B3.19% Conservative: 43 Query Match: 6 Gaps: US-10-622-516-2 (1-478) x I65510 (1-1824) Qy 1 MetProGlySerLeuProLeuAsnAlaGluAlaCysTrpProLysAspValGlyIleVal 20	TITLE JOURNAL ATURES BOUTC	equence 1 from patent US 5668001. 65510 65510.1 GI:2482080 nknown. nknown. nclassified. (bases 1 to 1824) liziorko, H.M.	477 GluHis 478 1610 GTGCAT 1615 SULT 15	Qy 438 GlyValGlyLeuValHisSerAsnIleAlaThrGluHisIleProSerProAlaLysLys 457	398 IleProGlnGlySerIleAspSerLeuPheGluGlyThrTrpTyrLeuValArgValA	90 ĠĊTĠĊTAĊĠĊTĠTATTĊĊATCAGAĠTTAĊAĊAĠGATĠĊCAĊTĊĊTĠĠTTĊTĠĊĠĊTTĠAĊ 124 58 LysIleThrAlaSerLeuCysAspLeuLysSerArgLeuAspSerArgThrGlyValAla 377	nAsnGlyAsnMetTyrThrSerSerValTyrGlySerLeuAlaSerValLeuAlaG
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                                The present invention relates to the isolation of a novel numan entry that shows a high degree of similarity to human cytoplasmic 3-hydroxy-3-12 methylglutaryl coenzyme A (HMG-CoA) synthase. The invention also discloses polynucleotide sequences encoding the novel enzyme of the invention. Both the polypeptide and polynucleotide sequences are useful as models for the development of human therapeutics, for identifying therapeutic proteins, as targets for development of human therapeutic agents, and as query sequences to perform a search against sequence capolypeptide is useful to raise antibodies or to elicit another immune response, as a reagent in assays designed to quantitatively determine levels of the protein in biological fluids, as markers for tissues in which the corresponding protein is preferentially expressed, in drug screening assays, in cell-based or cell-free systems, to identify compounds that modulate synthase activity of the protein in its natural state, or an altered form that causes the specific disease or pathology associated with the synthase, to screen a compound for the ability to stimulate or inhibit interaction between the synthase protein and a molecule that normally interacts with the synthase protein, and in
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WPI; 2(P-PSDB 2003-313241/30.)B; ABR54268.

Novel human proteins and nucleic acid encoding the proteins, useful for diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.

Claim 20; Page 245-246; 460pp; English.

The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiidiabetic, antiinfertility, haemostatic, netinflammatory, anti-HIV, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, valve disease, tuberous sclerosis, sclerosis, eclerosis, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, latentopoletic disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, Parkinson's disease, immune disorders, haemophilia, haemophilia, cancer, and metabolic syndrome x. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention. The human NOV35b protein in the exemplification of the present invention with the human NOV35b protein in the exemplification of the present invention.

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Human NOV41n encoding cDNA SEQ ID NO:207.

Human; NOVX; antiatherosclerotic; hypotensive; cardiant; dermatological; Wanorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; Wanorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; Waneuroprotective; nootropic; antiasthmatic; anti-HIV; immunomodulator; Waneuroprotective; nootropic; antiaparkinsonian; metabolic; antilipaemic; Waneuroprotective; nootropic; antiparkinsonian; metabolic; antilipaemic; Walve disease; transplantation; Walve disease; transplantation; Walve disease; transplantation; Walve disease; transplantation; Walve disease; metabolic disorder; neoplasm; lymphoma; uterus cancer; Walve disease; metabolic disorder; neoplasm; lymphoma; uterus cancer; Walve disease; multiple sclerosis; bronchial asthma; anorexia; Walve disease; multiple sclerosis; infectious disease; cancer; Walve disease; multiple sclerosis; infectious disease; cancer; Walve disease; haematopoietic disorder; dyslipidaemia; Wanetabolic syndrome X; gene; ss.

Homo sapiens

WO2 003023001-A2

0 MAR-2003

9 SEP-2002; 2002WO-US028538

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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in 2C hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, 2C antidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, 2C antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, 2C therapy. NOVX proteins are useful for treating or preventing a pathology 2C associated with a NOVX protein in humans and for treating a syndrome 2C antibodies can be used in the treatment and diagnosis of cardiomyopathy, 2C antivodies can be used in the treatment and diagnosis of cardiomyopathy, 2C antivodies can be used in the treatment and diagnosis of cardiomyopathy, 2C congenital adrenal hypertension, congenital heart defects, acrtic stenosis, 2C valve disease, tuberous sclerosis, scleroderma, obesity, transplantation, 2C disease, neoplasm, lymphoma, uterus cancer, diabetes, metabolic 2C disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, 2C hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host 2C disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, 2C Alzheimer's disease, parkinson's disease, immune disorders, 2C Alzheimer's disease, parkinson's disease, immune disorders, 2C Alzheimer's disease, anorexia, cancer-associated cachexia, cancer, 2C Algority and 2C Algorit
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Novel human proteins and nucleic acid encoding the proteins, useful for diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.
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Agee ML, Alsobrook JP, Burgess CE, Casman SJ, Crabtree J, Dipippo VA, Gangolli EA, Gerlach VL, Kekuda R, Khramtsov NV, Malyankar UM, Miller CE, Pena CEA, Rieger DK, Rc Spaderna SK, Spytek KA, Zerhusen BD, Zhong M; Rieger DK, Ro K, Spytek KA, D, Zhong M; Khramtsov NV, JM, Miller CE, Dipippo VA, Gerlach VL, CE, Ooi CE, Ort Rothenberg ME, KA, Taupier RJ, Giot L, Gorm
Leach MD, Lep Lepley DM, Li L Ort T, Padigaru I , Shenoy SG, Sh , Twomlow N, Ve Juo X, Gusev VY, Ji W;
Li L, Liu X;
Igaru M, Patturajan M;
3, Shimkets RA;
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Novel human diagnosis, t protein or n proteins and nucleic acid encoding the proteins, use treatment and prevention of disorders involving the hucleic acid e.g. cardiac and neurological disorders. useful for human

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ABR54167 to ABR54276. Nova hypotensive, dermatological, anorectic, immunocyte antidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-..., antiasthmatic, metabolic, immunomodulator, neuroprotective, nocropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, valve disease, tuberous sclerosis, scleroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, AIDS, bronchial asthma, Crohm's disease, multiple sclerosis, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention. The 1 the ant the ass ass ass ant the the hypological final hypological hypolo ent invention describes isolated human NOVX proteins, ACC62236 to ACC62345 encode the human NOVX proteins go to ABR54276. NOVX sequences have antiatheroscierotic, given in

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SULT 6 C62336 ACC62336 standard; cDNA; 1650

ACC62336;

23-JUN-2003

(first entry)

NOV41k encoding cDNA SEQ H NO:201.

Human; NOVX; antiatherosclerotic; hypotensive; cardiant; dermatological; anorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; kW haemostatic; antiinfelammatory; antiasthmatic; anti-HIV; immunomodulator; kW neuroprotective; nootropic; antiparkinsonian; metabolic; antilipaemic; kW congenital heart defect; aortic stenosis; hypertension; scleroderma; kW congenital heart defect; aortic stenosis; valve disease; transplantation; kW tuberous sclerosis; obesity; congenital adrenal hyperplasia; diabetes; kW prostate cancer; metabolic disorder; neoplasm; lymphoma; uterus cancer; kW idiopathic thrombocytopenic purpura; AIDS; bronchial asthma; anorexia; kW cancer-associated cachexia; Alzheimer's disease; cancer; cancer; haematopoietic disorder; dyslipidaemia; wetabolic syndrome X; gene; ss.

sapiens

WO2003023001-

20-MAR-2003

09-SEP-2002; 2002WO-US028538

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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host infectious disease, anorexia, cancer-associated cachexia, cancer, Alzhelmer's disease, parkinson's disease, immune disorders, cancer, caccer, which are used in examples from the present invention.

C ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention.
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Gangolli EA, Gerlach VL, Giot L, Gorman L, Guo X, Gusev VY, Ji W
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Pena CEA, Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA;
Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss E:
Zerhusen BD, Zhong M; Voss EZ; Σ.

WPI; DB; 2003-313241/30.)B; ABR54259.

Novel human proteins and nucleic acid encoding the proteins, useful diagnosis, treatment and prevention of disorders involving the human protein or nucleic acid e.g. cardiac and neurological disorders.

Claim 20; Page 241; 460pp; English.

The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in 2C ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antidabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiparkinsonian and antilipaemic activities, and can be used in gene cherapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, diabetes, metabolic disease, AIDS, bronchial asthma, Crohn's disease, mutiple sclerosis, cancer, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, AIDS, bronchial asthma, Crohn's disease, mutiple sclerosis, chematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention. ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention.

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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiatiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiaparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host cdisease, AlDS, bronchial asthma, Crohn's disease, mutiple sclerosis, ancer, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention. ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention.
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Burgess CE, Casman SJ, Catterton E, Chant JS, Chaudhuri A;
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Human; NOVX; antiatherosclerotic; hypotensive; cardiant; dermatological; W anorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; W haemostatic; antiinflammatory; antiasthmatic; anti-HIV; immunomodulator; W neuroprotective; nootropic; antiparkinsonian; metabolic; antilipaemic; W congenital heart defect; aortic stenosis; valve disease; transplantation; W tuberous sclerosis; obesity; congenital adrenal hyperplasia; diabetes; W prostate cancer; metabolic disorder; neoplasm; lymphoma; uterus cancer; W fertility; haemophilia; hypercoagulation; graft versus host disease; W idiopathic thrombocytopenic purpura; AIDS; bronchial asthma; anorexia; W cancer-associated cachexia; Alzheimer's disease; cancer; cancer; haematopoietic disorder; dyslipidaemia; W immune disorder; haematopoietic disorder; dyslipidaemia; Human -JUN-2003 NOV41e encoding ret CDNA S E Q H NO:18 Ó

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Ħ 20; Page 242; 460pp; English.

The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, Chypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antiasthmatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene therapy. NOVX proteins are useful for treating or preventing a pathology associated with a NOVX protein in humans and for treating a syndrome associated with the human disease. NOVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, cangenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercosquilation, idiopathic thrombocytopenic purpura, graft versus host cisease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, ancreations disease, anorexia, cancer-associated cachexia, cancer, ALDS, bronchial asthma, Crohn's disease, multiple sclerosis, chaematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent FCR primers and probes for human NOVX sequences, which are used in examples from the present invention.

Chaesaclastic antilipaemic activities, and metabolic syndrome X. ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention. quences, wnic... R54277 represents a

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ACC62338

23-JUN-2003 (first entry)

NOV41m encoding cDNA SEQ ID NO:205.

Human; NOVX; antiatherosclerotic; hypotensive; cardiant; dermatological; anorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; haemostatic; antiinflammatory; antiasthmatic; anti-HIV; immunomodulator; neuroprotective; nootropic; antiparkinsonian; metabolic; antilipaemic; gene therapy; cardiomyopathy; atherosclerosis; hypertension; scleroderma; congenital heart defect; aortic stenosis; valve disease; transplantation; tuberous sclerosis; obesity; congenital adrenal hyperplasia; diabetes; prostate cancer; metabolic disorder; neoplasm; lymphoma; uterus cancer; fertility; haemophilia; hypercoagulation; graft versus host disease; idiopathic thrombocytopenic purpura; AIDS; bronchial asthma; anorexia; cancer-associated cachexia; Alzheimer's disease; cancer; immune disorder; haematopoietic disorder; dyslipidaemia; metabolic syndrome X; gene; ss.

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2001US-0318120P.
2001US-0318184P.
2001US-0318430P.
2001US-0322636P.
2001US-0322816P.
2001US-0322817P.
2001US-0323519P.
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2001US-0323636P.
2001US-0324969P.
2001US-0325091P.
2001US-0325091P.
2001US-0341144P.
2002US-0361663P.
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Agee ML, Alsobrook JP, Anderson DW, Berghs C, Boldog FL;
Burgess CE, Casman SJ, Catterton E, Chant JS, Chaudhuri A;
Crabtree J, Dipippo VA, Edinger SR, Eisen AJ, Ellerman K;
Gangolli EA, Gerlach VL, Giot L, Gorman L, Guo X, Gusev VY, Ji W,
Kekuda R, Khramtsov NV, Leach MD, Lepley DM, Li L, Liu X;
Malyankar UM, Miller CE, Ooi CE, Ort T, Padigaru M, Patturajan M;
Pena CEA, Rieger DK, Rothenberg ME, Shenoy SG, Shimkets RA;
Spaderna SK, Spytek KA, Taupier RJ, Twomlow N, Vernet CAM, Voss E;
Zerhusen BD, Zhong M;

Voss EZ;

Boldog FL; Chaudhuri A; Ellerman K; Gugev VY,

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WPI; 2003-313241/30. P-PSDB; ABR54265.

human

RESULT 11
ACC62334
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Human; NOVX; antiatherosclerotic; hypotensive; cardiant; dermatological; anorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; haemostatic; antiinflammatory; antiasthmatic; anti-HIV; immunomodulator; neuroprotective; nootropic; antiparkinsonian; metabolic; antilipaemic; gene therapy; cardiomyopathy; atherosclerosis; hypertension; scleroderma; congenital heart defect; aortic stenosis; valve disease; transplantation; tuberous sclerosis; obesity; congenital adrenal hyperplasia; diabetes; prostate cancer; metabolic disorder; neoplasm; lymphoma; uterus cancer; fertility; haemophilia; hypercoagulation; graft versus host disease; idiopathic thrombocytopenic purpura; AIDS; bronchial asthma; anorexia; cancer-associated cachexia; Alzheimer's disease; parkinson's disease; immune disorder; haematopoietic disorder; dyslipidaemia; une disorder; abolic syndrom syndrome gene;

The present invention describes isolated human NOVX proteins, where X is CC 1 to 42. ACC62236 to ACC62345 encode the human NOVX proteins given in CC ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, Chypotensive, dermatological, ancorectic, immunosuppressive, cytostatic, antiidiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, CC antigathmatic, metabolic, immunomodulator, neuroprotective, nootropic, antiparkinsonian and antilipaemic activities, and can be used in gene CC therapy. NOVX proteins are useful for treating or preventing a pathology CC associated with a NOVX protein in humans and for treating a syndrome CC atherosclerosis, hypertension, congenital heart defects, aortic stenosis, CC valve disease, tuberous sclerosis, sclerosis, sclerosis, obesity, transplantation, CC congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, AIDS, bronchial asthma, Crohn's disease, metabolic disease, anorexia, cancer-associated cachexia, cancer, alzheimer's disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, Parkinson's disease, immune disorders, cancer, caccastic sequences, which are used in examples from the present invention with the human NOV35b protein in the exemplification of the present invention.

CC ABR54277 represents a human trypsinogen protein given in comparison with the human NOV35b protein in the exemplification of the present invention. Novel human proteins and nucleic acid encoding the proteins, use diagnosis, treatment and prevention of disorders involving the protein or nucleic acid e.g. cardiac and neurological disorders. sequences, which are an ABR54277 represents a human human NOV35b protein in 20; Page 244; 460pp; English.

Sequence 1650 B₽; 477 A 319 397 G; 0 0 Other;

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ACC62327;

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Hum an NOV41b encoding cDNA SEQ ID NO:183

Human; NOVX; antiatherosclerotic; hypotensive; cardiant; dermatological; anorectic; immunosuppressive; cytostatic; antidiabetic; antiinfertility; haemostatic; antiinflammatory; antiasthmatic; anti-HIV; immunomodulator; neuroprotective; nootropic; antiparkinsonian; metabolic; antilipaemic; gene therapy; cardiomyopathy; atherosclerosis; hypertension; scleroderma; congenital heart defect; aortic stenosis; valve disease; transplantation; tuberous sclerosis; obesity; congenital adrenal hyperplasia; diabetes; prostate cancer; metabolic disorder; neoplasm; lymphoma; uterus cancer; fertility; haemophilia; hypercoagulation; graft versus host disease; idiopathic thrombocytopenic purpura; AIDS; bronchial asthma; anorexia; Crohn's disease; multiple sclerosis; infectious disease; cancer; cancer; haematopoietic disorder; disease; Parkinson's disease; immune disorder; haematopoietic disorder; dyslipidaemia;

Hom Ō sapiens

WO2 003023001-A2

0 MAR-2003

ø SEP-2002; 2002WO-US028538.

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The present invention describes isolated human NOVX proteins, where X is 1 to 42. ACC6236 to ACC62345 encode the human NOVX proteins given in ABR54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant, hypotensive, dermatological, anorectic, immunosuppressive, cytostatic, antiadiabetic, antiinfertility, haemostatic, antiinflammatory, anti-HIV, antisathmatic, metabolic, immunomodulator, neuroprotective, nootropic, associated with an NOVX protein are useful for treating or preventing a pathology associated with the human disease. NOVX nucleic acids, proteins and easpeciated with the human disease. NOVX nucleic acids, proteins and atherosclerosis, hypertension, congenital heart defects, aortic stenosis, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disease, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, anorexia, cancer-associated cachexia, cancer, Alzheimer's disease, anorexia, cancer-associated cachexia, cancer, haematopoletic disorders, dyslipidaemias, and metabolic syndrome X. ACC623465 represent PCR primers and metabolic syndrome X. Charles which are used in examples from the present invention. The human NOV35b protein in the exemplification of the present invention.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Agee ML, Alsobrook JP, Anderson DW, IBurgess CE, Casman SJ, Catterton E, Crabtree J, Dipippo VA, Edinger SR, IGangolli EA, Gerlach VL, Giot L, Gorr Kekuda R, Khramtsov NV, Leach MD, Leimalyankar UM, Miller CE, Ooi CE, Ort Pena CEA, Rieger DK, Rothenberg ME, Spaderna SK, Spytek KA, Taupier RJ, Zerhusen BD, Zhong M;
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17-SEP-2001;

17-SEP-2001;

19-SEP-2001;

20-SEP-2001;

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25-SEP-2001;

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2001US-0324969P.
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Nove diag ignosis, itein or el human n proteins and nucleic acid encoding the proteins, useful treatment and prevention of disorders involving the human nucleic acid e.g. cardiac and neurological disorders. useful for

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ABR54167 to ABR54276. NOVX sequences

Appotensive, dermatcological, anorectic, immunosuppressive, original properties, immunosuppressive, anti-HIV, antidabetic, antiinfertility, haemostatic, neuroprotective, nootropic, antiparkinsonian and antilpaemic activities, and can be used in gene therapy. NovX proteins are useful for treating or preventing a pathology associated with a NoVX protein in humans and for treating a syndrome associated with the human disease. NoVX nucleic acids, proteins and antibodies can be used in the treatment and diagnosis of cardiomyopathy, atherosclerosis, hypertension, congenital heart defects, aortic stenosis, valve disease, tuberous sclerosis, scleroderma, obesity, transplantation, congenital adrenal hyperplasia, prostate cancer, diabetes, metabolic disorders, neoplasm, lymphoma, uterus cancer, fertility, haemophilia, hypercoagulation, idiopathic thrombocytopenic purpura, graft versus host disease, AIDS, bronchial asthma, Crohn's disease, multiple sclerosis, largeticus disease, Parkinson's disease, immune disorders, cancer, Alzheimer's disease, Parkinson's disease, immune disorders, cancer, haematopoietic disorders, dyslipidaemias, and metabolic syndrome X. ACC62346 to ACC62465 represent PCR primers and probes for human NOVX sequences, which are used in examples from the present invention.

The examples from the present invention with processin invention. The 1 to ABRI ABRI antiantiantiantiantiantiantithe association the association the association the association that antian present invention describes isolated human NOVX proteins, where X is o 42. ACC62236 to ACC62345 encode the human NOVX proteins given in 54167 to ABR54276. NOVX sequences have antiatherosclerotic, cardiant,

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                                                                                                                                                                             This invention relates to a novel method of treating a human subject having a tumourigenic disorder or angiogenic disorder, caused by aberrant gene expression or activity of an isolated protein, by administering a modulator. The modulator may have cytostatic, antithyroid, antidiabetic or ophthalmological activity. The method is useful for treating a subject having a tumourigenic or angiogenic disorder, in particular for treating cancer (for example breast cancer, colon cancer, lung cancer or prostatic cancer) and, for example, Grave's disease and diabetic retinopathy. The present sequence is a DNA sequence which encodes the novel isolated human protein 9389 of the invention.
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Qy	319 SerProGlnGlnLeuAlaGlyLysArgIleGlyValPheSerTyrGlySerGlyLeuAla 338
da	1329 TCACCTCAGCAATTAGCAGGGAAGAGAATTGGAGTGTTTTCTTATGGTTCTGGTTTGGCT 1388
γQ	339 AlaThrLeuTyrSerLeuLysValThrGlnAspAlaThrProGlySerAlaLeuAspLys 358
đđ	1389 GCCACTCTGTACTCTCTTAAAGTCACACAAGATGCTACACCGGGGTCTGCTCTTGATAAA 1448
Qy	359 IleThrAlaSerLeuCysAspLeuLysSerArgLeuAspSerArgThrGlyValAlaPro 378
da	1449 ATAACAGCAAGTTTATGTGATCTTAAATCAAGGCTTGATTCAAGAACTGGTGTGGCACCA 1508
ઇ	379 AspValPheAlaGluAsnMetLysLeuArgGluAspThrHisHisLeuValAsnTyrIle 398
dd	1509 GATGTCTTCGCTGAAAACATGAAGCTCAGAGAGGACACCCATCATTTGGTCAACTATATT 1568
8	399 ProGlnGlySerIleAspSerLeuPheGluGlyThrTrpTyrLeuValArgValAspGlu 418
d d	1569 CCCCAGGGTTCAATAGATTCACTCTTTGAAGGAACGTGGTACTTAGTTAG
γO	419 LysHisArgArgThrTyrAlaArgArgProThrProAsnAspAspThrLeuAspGluGly 438
Db	1629 AAGCACAGAAGAACTTACGCTCGGCGTCCCACTCCAAATGATGACACTTTGGATGAAGGA 1688
Ş	439 ValGlyLeuValHisSerAsnIleAlaThrGluHisIleProSerProAlaLysLysVal 458
ממ	1689 GTAGGACTTGTGCATTCAAACATAGCAACTGAGCATATTCCAAGCCCTGCCAAGAAAAGTA 1748
Qy	459 ProArgLeuProAlaThrAlaAlaGluProGluAlaAlaValIleSerAsnGlyGluHis 478
дb	1749 CCAAGACTCCCTGCCACAGCAGCAGAACCTGAAGCAGCTGTCATTAGTAATGGGGAACAT 1808
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### Pred. No.: Pred. No.: 2.39e-293	CURRENT FILING DATE: 2003-07-21 PRIOR APPLICATION NUMBER: 10/193,295 PRIOR FILING DATE: 2002-07-12 PRIOR APPLICATION NUMBER: 09/819,993 PRIOR FILING DATE: 2001-03-29 NUMBER OF SEQ ID NOS: 5 SOFTWARE: FastSEQ for Windows Version 4. SEQ ID NO 1 LENGTH: 2002 TYPE: DNA ORGANISM: Human S-10-622-516-1	22-516-1 22-516-1 ace 1, Application US/10622516 cation No. US20040018545A1 at information: ICANT: GONG, Fangcheng et al. ICANT: GONG, Fangcheng et al. ICANT: OF INVENTION: ISOLATED HUMAN ENZYME PROTEINS, NUCLEIC OF INVENTION: ACID MOLECULES ENCODING HUMAN ENZYME PROTEINS OF INVENTION: THEREOF INVENTION: THEREOF REFERENCE: CL001195DIV2 ENT APPLICATION NUMBER: US/10/622,516	Qy 421 ArgArgThrTyrAlaArgArgProThrProAsnAspAspThrLeuAspGluGlyValGly 440	Qy 321 GlnGlnLeuAlaGlyLysArgIleGlyValPheSerTyrGlySerGlyLeuAlaAlaThr 340

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	TYPE: DNA ORGANISM: Homo sapiens FEATURE: NAME/KEY: CDS LOCATION: (2)(1562) S-10-236-417-207 lignment Scores:	PRIOR FILING DATE: 2002-07-17 PRIOR APPLICATION NUMBER: US60/396,412 PRIOR FILING DATE: 2002-07-17 PRIOR APPLICATION NUMBER: US60/322,636 PRIOR FILING DATE: 2001-09-17 PRIOR FILING DATE: 2001-09-17 PRIOR APPLICATION NUMBER: US60/322,816 PRIOR APPLICATION NUMBER: US60/322,816 PRIOR APPLICATION NUMBER: US60/323,519 PRIOR APPLICATION NUMBER: US60/323,519 PRIOR FILING DATE: 2001-09-19 Remaining Prior Application data removed - See File NUMBER OF SEQ ID NOS: 341 SOFTWARE: Custom SEQ ID NO 207 LENGTH: 1564	Sequence 207, Application US/10236417 Publication No. US20040048256A1 GENERAL INFORMATION: APPLICANT: Agee et al. TITLE OF INVENTION: NOVEL PROTEINS AND NUCLEIC ACIDS FILE REFERENCE: 21402-442C CURRENT APPLICATION NUMBER: US/10/236,417 CURRENT FILING DATE: 2003-01-06 PRIOR APPLICATION NUMBER: US60/318,120 PRIOR APPLICATION NUMBER: US60/318,430 PRIOR FILING DATE: 2001-09-01 PRIOR APPLICATION NUMBER: US60/318,430 PRIOR APPLICATION NUMBER: US60/322,781 PRIOR APPLICATION NUMBER: US60/322,781 PRIOR APPLICATION NUMBER: US60/322,781 PRIOR APPLICATION NUMBER: US60/320,781 PRIOR APPLICATION NUMBER: US60/318,184 PRIOR APPLICATION NUMBER: US60/318,184 PRIOR APPLICATION NUMBER: US60/361,663	1142 GCCACTCTGTACTCT 359 IleThrAlaSerLeu

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Sequence 369, Application US/10307817;

Publication No. US20040058338A1;

GENERAL INFORMATION:

APPLICANT: Agee et al.

TITLE OF INVENTION: NOVEL PROTEINS AND NOTE OF ILE REFERENCE: 21402-502C;

CURRENT APPLICATION NUMBER: US/10/307,81.

CURRENT FILING DATE: 2002-12-02;

NUMBER OF SEQ ID NOS: 682;

SOFTWARE: CuraSeqList version 0.1;

SEQ ID NO 369;

LENGTH: 1564;

TYPE: DNA

ORGANISM: Homo sapiens
FEATURE:

NAME/KEY: CDS
LOCATION: (2)..(1561)

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; Publication No. US20040058338A1
; GENERAL INFORMATION:
; APPLICANT: Agee et al.
; TITLE OF INVENTION: NOVEL PROTEINS AND NUCLE
; FILE REFERENCE: 21402-502C
; CURRENT APPLICATION NUMBER: US/10/307,817
; CURRENT APPLICATION NUMBER: US/10/307,817
; CURRENT FILING DATE: 2002-12-02
; NUMBER OF SEQ ID NOS: 682
; SOFTWARE: CuraSeqList version 0.1
; SEQ ID NO 373
; LENGTH: 1564
; TYPE: DNA
; ORGANISM: Homo sapiens
; PEATURE:
; NAME/KEY: CDS
; LOCATION: (2)..(1561)
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RIOR APPLICATION NUMBER: US60/318,120

RIOR FILING DATE: 2001-09-01

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R APPLICATION NUMBER: US60/318,430

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R FILING DATE: 2001-09-17

R APPLICATION NUMBER: US60/322,781

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R APPLICATION NUMBER: US60/318,184

R APPLICATION NUMBER: US60/361,663

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R APPLICATION NUMBER: US60/396,412

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Sequence 189, Application US/10236417

publication No. US20040048256A1

GENERAL INFORMATION:
APPLICANT: Agee et al.

FITTLE OF INVENTION: NOVEL PROTEINS AND NUCLE
FILE REFERENCE: 21402-442C

CURRENT APPLICATION NUMBER: US/10/236,417

CURRENT FILING DATE: 2003-01-06

PRIOR APPLICATION NUMBER: US60/318,120

PRIOR FILING DATE: 2001-09-01

PRIOR APPLICATION NUMBER: US60/318,430

PRIOR FILING DATE: 2001-09-17

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PRIOR APPLICATION NUMBER: US60/322,636

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PRIOR APPLICATION NUMBER: US60/322,636

PRIOR FILING DATE: 2001-09-17

PRIOR APPLICATION NUMBER: US60/322,816

PRIOR APPLICATION NUMBER: US60/323,519

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US-10-236-417-193

US-10-236-417-193

; Sequence 193, Application US/10236417
; Publication No. US20040048256A1
; GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND NUCLE
FILE REFERENCE: 21402-442C
CURRENT APPLICATION NUMBER: US/10/236,417
; CURRENT FILING DATE: 2003-01-06
; PRIOR APPLICATION NUMBER: US60/318,120
pRIOR APPLICATION NUMBER: US60/318,430
pRIOR APPLICATION NUMBER: US60/318,430
pRIOR APPLICATION NUMBER: US60/322,781
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pRIOR APPLICATION NUMBER: US60/318,184
pRIOR FILING DATE: 2001-09-07
; PRIOR APPLICATION NUMBER: US60/361,663
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PRIOR FILING DATE: 2002-07-17
PRIOR APPLICATION NUMBER: US60/322,63;
PRIOR FILING DATE: 2001-09-17
PRIOR APPLICATION NUMBER: US60/322,81;
PRIOR APPLICATION NUMBER: US60/322,81;
PRIOR FILING DATE: 2001-09-17
PRIOR FILING DATE: 2001-09-17
PRIOR FILING DATE: 2001-09-19
Remaining Prior Application data removement of SEQ ID NOS: 341
SOFTWARE: Custom
SEQ ID NO 193
LENGTH: 1650
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: CDS
LOCATION: (22)..(1582)
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; Sequence 197, Application US/1023641
; Publication No. US20040048256A1
; GENERAL INFORMATION:
; APPLICANT: Agee et al.
; TITLE OF INVENTION: NOVEL PROTEINS
; FILE REFERENCE: 21402-442C
; CURRENT APPLICATION NUMBER: US/10/2
; CURRENT FILING DATE: 2003-01-06
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PRIOR FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,43;
PRIOR FILING DATE: 2001-09-10
PRIOR APPLICATION NUMBER: US60/322,78;
PRIOR APPLICATION NUMBER: US60/322,78;
PRIOR APPLICATION NUMBER: US60/361,66;
PRIOR APPLICATION NUMBER: US60/361,66;
PRIOR APPLICATION NUMBER: US60/396,41;
PRIOR APPLICATION NUMBER: US60/322,63;
PRIOR APPLICATION NUMBER: US60/322,63;
PRIOR APPLICATION NUMBER: US60/322,81;
PRIOR FILING DATE: 2001-09-17;
PRIOR APPLICATION NUMBER: US60/322,81;
PRIOR FILING DATE: 2001-09-17;
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PRIOR APPLICATION NUMBER: US60/322,81;
PRIOR FILING DATE: 2001-09-17;
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(Sequence 201, Application US/10236417
) Publication No. US20040048256A1

(GENERAL INFORMATION:
APPLICANT: Agee et al.
TITLE OF INVENTION: NOVEL PROTEINS AND NUCLE
FILE REFERENCE: 21402-442C
CURRENT APPLICATION NUMBER: US/10/236,417
CURRENT FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,120
PRIOR FILING DATE: 2001-09-01
PRIOR APPLICATION NUMBER: US60/318,430
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/322,781
PRIOR APPLICATION NUMBER: US60/361,663
PRIOR FILING DATE: 2001-09-07
PRIOR APPLICATION NUMBER: US60/361,663
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-Q=/cgn2_1/USPTO_spool_p/US10622516/runat_23062004_162634_673/app_query.fasta
-DB=Issued Patents_NA -QFMT=fastap -SUFFIX=rni -MINMATCH=0.1 -LOOPCL=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOCALIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=pto -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000
-USER=US10622516_@CGN_1_1_105_@runat_23062004_162634_673 -NCPU=6 -ICPU=3
-NO_MMAP -LARGEQUERY -NEG_SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=7
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Sequence 1, Application US/08305505

Patent No. 5668001

GENERAL INFORMATION: 3-HYDROXY-3-METHY

TITLE OF INVENTION: 3-HYDROXY-3-METHY

TITLE OF INVENTION: SYNTHASE PREPARAT

CORRESSES: Quarles & Brady

STREET: 411 East Wisconsin Avenue

CITY: Milwaukee

STATE: Wisconsin

COUNTRY: U.S.A.

ZIP: 53202

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER READABLE FORM:

COMPUTER: US.A.

ZIP: 5202

COMPUTER READABLE FORM:

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Mismatches:
Indels:
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; Patent No. 6436692
; GENERAL INFORMATION:
; APPLICANT: GONG, Fangcheng et al.
TITLE OF INVENTION: ISOLATED HUMAN ENZ)
; TITLE OF INVENTION: ACID MOLECULES ENC
TITLE OF INVENTION: THEREOF
; FILE REFERENCE: CL001195
; CURRENT APPLICATION NUMBER: US/09/819,9
; NUMBER OF SEQ ID NOS: 5
; SOFTWARE: FastSEQ for Windows Version 4
; SEQ ID NO 3
; LENGTH: 28001
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
NAME/KEY: misc_feature
; LOCATION: (1)...(28001)
; OTHER INFORMATION: n = A,T,C or G
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60	Db 19144 TTGGACTACAGCTGTGATTCAGGGAAAGCTAATGAAAATGAATTACTAAAGTGATCTTAC 19203 Qy
33 C3ACAATTTACACACACACACCTTACTTAACACACACAAAAAA	Qy 204 204
60	Db 19084 GGAGAAAAGAAGGTAAGTTGAATTTTGTTCATCTTTTGAGAGAGGGTATTTTAACAAGGTT 19143
	Qy 204 204
03 IIICCIICIAIAIIAGAGCAIIIIAAIAICIGIIAAGCIGIIAIIIGIACAGACCIGAG 2010	Db 19024 CTGAGGTGTATCTCTGCAAAAATATTTAGGTCGGTTTACCCCCTTGTAAGAAAATCAAAGT 19083
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44 TAAAATAGTATCTATAGTGGCCTGGAAGCCTTTGG-GTAAGAGGAGCTATTATGAGTTT 50	Db 18964 ACAACATTTTCCCATAGTTTCTGGGAAAGTGTAATTTACTAGAAGAGGTAAACTTTGGAA 19023
48 pLysAsnSerIleTyrSerGlyLeuGluAlaPheGly	204
19984 GGTTCAGAAATCTCTAGCTCGGATGTTGCTGAATGACTTCCTTAATGACCAGAATAGAGA 20043	Db 18904 AACTACTTTGTGGGCATTCTTCATTTAATATCCTTTTACCATTAATTCCTCATTCACCAA 18963
alGlnLysSerLeuAlaArgMetLeuLeuAsnAspPheLeuAsnAspG ;	18844 TTTTCCTTGGTTTTGGTATGAGTTGAGAGCAGTCTAATGTACTAGGTATCTTTGGTAGGC
208 pLysAspPheThrLeuAsnAspPheGlyPheMetIlePheHisSerProTyrCysLysLe 228	204
	Qy 190 CysTyrSerValTyrCysLysLysIleHisAlaGlnTrpGln-Lys 204
19804 TTGTGGTTGCTGTTTAAACATATACAAATATCCTAGCTTTATTCTAAAGTCAAACTTTAA 19863 205GluGlvAgnAs 208	Db 18724 GAATATCCTATAGTAGATGGAAAACTCTCCATACAGTGCTACCTCAGTGCATTAGACCGC 18783 Ov
	TyrProileValAspGlyLysLeuSerileGlnCysTyrLeuSerAlaLeuAspArg 189
19744 TTTGAGTGTCTTTTTAAATGTATACTTTAAGGTATAGAGAGGGTTTCATTATACAGTGTAT 19803	1866
204 204	\(1-4/8\) X US-10-193-295-3 \(1-28001\)
19684 CTGCTAATGTGTATGAATCTTAAATTTGAAAATTAGTGACATAGTACATATTGTTTCATC 19743	Gaps:
204 204	est Local Similarity: 19.16% Mismatches: uery Match: 33.43% Indels:
19624 TAAAATACTCTTTTATTTAAAAAAAATACTAATCCTGACCCACTAAATTGATTATGTAAC 19683	rcent Similarity: 19.23% Conservations
204 204	red. No.: 5.76e-92 Length:
19564 CATTACTTACCCTTGCTGCAAGTTATTCAGTTTGCTATTTTTCTACTGCATTTTGTTTTT 19623	S-10-193-295-3
204 204	OTHER INFORMATION: n
19504 ACAGTAAGAAGTACATGTTACATTGTATGTGTATGCCAGACTGAAACAAAAATGTCATGA 19563	NAME/KEY: misc_feature
204 204	ORGANIS
19444 AAGCATTCTTGCAGTATATTAACAGAATAGTGGTTTTCTAACTTTTTTATTAGGACCC 19503	C.
204 204	SOFTWARE: FastSEQ f
19384 GAAGGGGTTAAAAATCATATTCAATGACAAATATCAGTGAATTTAGTCGCTCTGGATAAG 19443	PRIOR FILING DATE: 2001-03-29
204 204	CURRENT FILING DATE: 2002-07-12 CURRENT FILING DATE: 2002-07-12
19324 CACATTATTACATAAAGTATACTTTTTCTGTAGTCCAACTTTGCTTTTTAGAGGTTATGA 19383	FILE REFERENCE: CLOOI195DIV
204 204	ACID
19264 ATTAATCTGGAAATTTGTTGAGGCACTGAAAGGACAGTATTTGAGTTAATGCTATCATAA 19323	APPLICANT: GONG, Fangcheng et al.
204 204	Sequence 3, Application Patent No. 6620608
19204 CCCAAAAATAATCTTTTTGCACTTGACCTGTGAATTTGTATTTGTTTTTTTACTGTTATC 19263	S-10-193-295-3
204 204	

22443 GGTGCAATGGCTCACACCTATAATCCCAACACTTCAGGAGGCTGAGGTGGGAGGATTGCT 22502	317 317	, 84 ,
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352 352	21003 AGATGTGGAGAAGGCATTTATGAAGGCTAGCTCTGAACTCTTCAGTCAG	ם מ
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	20943 TGTAAAACTGTAAAATAAATCTTTCAGGGATGTTAAATTAGAAGACACCTACTTTGATAG 21002	ДЬ
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RESULT 6
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; Sequence 245, Application US/093;
Patent No. 6444425
; Patent No. 6444425
; GENERAL INFORMATION:
; APPLICANT: Reed, Steven G.
; APPLICANT: Lodes, Michael J.
APPLICANT: Mohamath, Roadoh
; APPLICANT: Secrist, Heather
; TITLE OF INVENTION: COMPOUNDS F
; TITLE OF INVENTION: LUNG CANCER
; FILE REFERENCE: 210121.475C1
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Y Ma
  PLICANT: Reed, Steven G.
PLICANT: Lodes, Michael J.
PLICANT: Mohamath, Roadoh
PLICANT: Secrist, Heather
PLICANT: Secrist, Heather
PLE OF INVENTION: COMPOUNDS FOR THERAF
PLE OF INVENTION: LUNG CANCER AND METH
LE REFERENCE: 210121.475C1
RENT APPLICATION NUMBER: US/09/370,83
RENT FILING DATE: 1999-08-09
RLIER APPLICATION NUMBER: US 09/285,32
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RENT FILING DATE: 1999-08-09
LIER APPLICATION NUMBER: US 09/285,3
LIER FILING DATE: 1999-04-02
BER OF SEQ ID NOS: 289
TWARE: FastSEQ for Windows Version 3
ID NO 245
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E/KEY: misc
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RESULT 8
US-09-401-064-187
; Sequence 187, Application US/0940;
patent No. 6623923
; GENERAL INFORMATION:
APPLICANT: Xu, Jiangchun
APPLICANT: Lodes, Michael J.
APPLICANT: Secrist, Heather
APPLICANT: Benson, Darin R.
APPLICANT: Meagher, Madeline Joy
APPLICANT: Wang, Tongtong
TITLE OF INVENTION: COMPOUNDS FOR TITLE OF INVENTION: DIAGNOSIS OF FILE REFERENCE: 210121.471C2; CURRENT APPLICATION NUMBER: US/CURRENT FILING DATE: 1999-09-22; NUMBER OF SEQ ID NOS: 371
SOFTWARE: FastSEQ for Windows Very SEQ ID NO 187
LENGTH: 506
TYPE: DNA
ORGANISM: Homo sapien
US-09-401-064-187
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NUMBER OF SEQ ID NOS: 289
SOFTWARE: FastSEQ for Windows V
SEQ ID NO 128
LENGTH: 500
TYPE: DNA
ORGANISM: Homo sapien
S-09-370-838-128
Alignment
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                             US-09-306-595C-1

; Sequence 1, Application US/09306595C

; Patent No. 6284506

; GENERAL INFORMATION:

; APPLICANT: HOSHINO, Tatsuo

APPLICANT: SITOGUCHI, Yutaka

APPLICANT: SITOGUCHI, Yutaka

TITLE OF INVENTION: ISOPRENIOD PRODUCTION

; FILE REFERENCE: ISOPRENIOD PRODUCTION

; CURRENT APPLICATION NUMBER: US/09/306,595C

CURRENT APPLICATION NUMBER: US/09/306,595C

CURRENT FILING DATE: 1998-05-06

PRIOR APPLICATION NUMBER: 98108210

PRIOR FILING DATE: 1998-05-06

NUMBER OF SEQ ID NOS: 43

SOFTWARE: PatentIN Ver. 2.1

; SEQ ID NO 1

LENGTH: 4775

TYPE: DNA

ORGANISM: Phaffia rhodozyma

FEATURE:

NAME/KEY: exon

LOCATION: (1305)...(1361)

NAME/KEY: intron

LOCATION: (1362)...

LOCATION: (1505)...(1504)

NAME/KEY: intron

LOCATION: (1505)...(1509)

NAME/KEY: exon

LOCATION: (1500)...(1826)
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TCTGTCAAGACAGTCCTTATGGACTTGTTCGAGTCCCACGG IleAspThrThrAsnAlaCysTyrGlyGlyThrAlaAlaVa	102	Qy 82 uSerTyrAspCysIleGlyArgLeuGluValGlyThrGluThrIleIleAspLysSerLy 102 : :::::	Qy 73	1832 TCTCTTCCGTTTCAGCAATCGACAGGAAAAAAGGCCCCAAGCGCATCTCACTGACACCTTTC	72	Qy 54 AlaLysMetGlyPheCysThrAspArgGluAspIleAsnSerLeuCysMet-ThrVal 72	Qy 35GluLeuGluLyBTyrAspGlyValAspAlaGlyLySTyrThrIleGlyLeuGlyGln 53 ::: ::: :::	:::::: 1652 TATCTCCCGACGCGAAATACAACACTGACCGCGATTTCTCTCGATCAGGCCATCGCTCAC	\rPheProSerGlnTyrVal	ery Match: 23.30% Indels: 3 Gaps: 3	1.36e-61 Length 585.00 Matche Similarity: 37.15% Conser al Similarity: 25.56% Mismat	LOCATION: (4043) -09-306-595C-1 ignment Scores:	LOCATION: (3602)(3768 NAME/KEY: polyA_site	LOCATION: (3:	NAME/KEY: exon LOCATION: (3326)(3493	NAME/KEY: intron LOCATION: (3241)(3325	LOCATION: (2892). NAME/KEY: exon	LOCATION: (2 NAME/KEY: in	LOCATION: (27	NAME/KEY: EXCH	LOCATION: (25	LOCATION: (249	LOCATION: (24	/KEY: exon TION: (2352)(2409	; LOCATION: (1921)(2277) ; NAME/KEY: intron ; IOCATION: (2278) (2351)	NAME/KEY: exo
287 nLysThrLysAlaSerLeuLeuValSerAsnGlnAsnGlyAsnMetTyr: :::: :::	Qy 267 rTyrPheAspArgAspValGluLysAlaPheMetLysAlaSerSerGluLeuPheSerGl 287 :	Qy 248 pLysAsnSerIleTyrSerGlyLeuGluAiaPheGlyAspVaiLysLeuGluAspTh 267 ::: :::::: Db 3016GACCCGGTTTTTGCTGAGGTGCCAGCCGAGCTTGCTACTTTGGACATGAAGAAAAG 3071	23/	2901 CTTTTTGTAACTCTTAGCTTGCAGATAAAACTTTTAGGTTTCTGGTACTCATTATTTAT	Qy 236 236	Qy 226 CysLysLeuValGlnLysSerLeuAlaArgMet 236		21 GGTTGCCGGTGTCAGTGCTGCGTCGTTCGATTACCTTTTGTTCCA	10AspPheThrLeuAsnAspPheGlyPheMetIlePheH :::::::::	Oy 199 sAlaGlnTrpGlnLysGluGlyAsnAspLys	Qy 179 rIleGlnCysTyrLeuSerAlaLeuAspArgCysTyrSerValTyrCysLysIleHi 199 ::: ::: ::: :::	Qy 170GluTyrProIleValAspGlyLysLeuSe 179		Qy 169 169	2431 TGCGCTTGGGAGAGTCTTACACTAATTCGGGGTGCTCGTATCCTTCGAATCGTTTGTTGC	69	156 tGlnHisAlaTyrAspPheTyrLysProAspMetLeuSer	Db 2311 GCGGCTGAAACCAACCCTTATCCGTCATTCTCATCAATCTAGCCGTCCACGGAAACTTCAT 2370	53	Db 2251 GACCCGACGCTCCCGTCGTCTTCGAGCGTGAGTTCCAATCCGTCATTTTCTTCCACGGCA 2310	Qy 152 152	Db 2191 TCTACGCCGAGGGTGCTGCCCGACCTGCCGGAGGTGCTGGTGCTTGCGCCATCCTCATCG 2250	Qy 152 152	Db 2132 GATCGAGTCATCCTCTTGGGACGGAAG-AAATGCCATTGTCTTCTGCGGAGACATTGCCA 2190	2072 TATCGACTCCAAGAATGCCTGCTACGGTTCTACCGCGGCCCTGTTCAATGCCGTCAACTG	

	RESULT 10 US-09-925-388-1 ; Sequence 1, Application US/09925388 ; Patent No. 6586202 ; GENERAL INFORMATION: ; APPLICANT: HOSHINO, Tatsuo APPLICANT: OJIMA, Kazuyuki APPLICANT: SETOGUCHI, Yutaka ; TITLE OF INVENTION: ISOPRENOID PRODUCTION ; FILE REFERENCE: ISOPRENOID PRODUCTION ; CURRENT APPLICATION NUMBER: US/09/925,388 ; CURRENT FILING DATE: 2001-08-09 ; PRIOR APPLICATION NUMBER: 09/306,595 ; PRIOR FILING DATE: 1999-05-06 ; NUMBER OF SEQ ID NOS: 43 ; SOFTWARE: PatentIn Ver. 2.1 ; EQ ID NO 1 ; LENGTH: 4775	353 353 365 365 371	3312 TTTCTCTGAACAGGTCGGCAAGCGCATTGCTCTACGCCTACGGATCTGGAGCTGCTG 3 339 laThrLeuTyrSerLeuLysValThrGlnAspAlaThrProGlySerAlaLeuAspLysI 3 :::	3132 GCAGGTTGAGCCTGGAATGACCACCGTCCGACAGCTCGGAAACTTGTACACCGCCTCTCT 319 307 lTyrGlySerLeuAlaSerValLeuAlaGlnTyrSerProGlnGlnLeu 323 ::: :: :: 3192 CTTCGGTGCTCTCGCAAGTTTTGTTCTCTAATGTTCCTGGTGACGAGCTCGTAAGTCTTGA 325 323
Db 1832 TCTCTTCCGTTTCAGCAATCGACAGGAAAAAGGCCCAAGCGCATCTCACTGACACCTTTC 1891 Qy 73	rcent Similarity: 37.15% Conservative: 82 st Local Similarity: 25.56% Mismatches: 12 ery Match: 4 23.30% Indels: 31 -10-622-516-2 (1-478) x US-09-925-388-1 (1-4775) 25 TyrPheProSerGlnTyrVal	ATION: (3241)(3325) E/KEY: exon ATION: (3326)(3493) E/KEY: intron ATION: (3494)(3601) E/KEY: exon E/KEY: exon ATION: (3602)(3768) E/KEY: polyA_site ATION: (4043)(4044) 25-388-1 nt Scores:	LOCATION: (2410)(2497 NAME/KEY: exon LOCATION: (2498)(2504 NAME/KEY: intron LOCATION: (2505)(2586 NAME/KEY: exon LOCATION: (2587)(2768 NAME/KEY: intron LOCATION: (2769)(2851 NAME/KEY: exon LOCATION: (2852)(2891 NAME/KEY: intron LOCATION: (2892)(2985 NAME/KEY: exon LOCATION: (2892)(2985 NAME/KEY: exon LOCATION: (2892)(2985 NAME/KEY: exon LOCATION: (2986)(3240	(1523)(1699 exon (1700)(1826 intron (1827)(1920 exon (1921)(2277 intron (2278)(2351 exon (2352)(2409)

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APPLICANT: Knechtle, Philipp APPLICANT: Rebischung, Corinne ITITLE OF INVENTION: GENOMIC DNA SEQUENCES OF ASHBYA GOSSYPII ITITLE OF INVENTION: AND USES THEREOF NUMBER OF SEQUENCES: 1152 CORRESPONDENCE ADDRESS: ADDRESSEE: No. 6239264artis Corporation STREET: 3054 Cornwallis Road CITY: Research Triangle Park STATE: No. 6239264th Carolina COUNTRY: USA IP: 27709 COMPUTER READABLE FORM: MEDIUM TYPE: Floppy disk COMPUTER: IBM PC compatible OPERATING SYSTEM: PC-DOS/MS-DOS	SULT 11 -08-998-416-730 Sequence 730, Application US/0899 Patent No. 6239264 GENERAL INFORMATION: APPLICANT: Philippsen, Peter APPLICANT: Pohlmann, Rainer APPLICANT: Steiner, Sabine APPLICANT: Mohr, Christine APPLICANT: Wendland, Jurgen	Qy 386 386 Db 3531 TTTGTCAACCGCTAACAACCTTCTTGAATCGGTCTCTTTTGAAATTCGCTCGGCG 3590 Qy 387LeuArgGluAspThrHisHisLeuValAsnTyrIleProGlnGlySerI 403 : : : :	Qy 339 laThrLeuTyrSerLeuLysValThrGlnAspAlaThrProGlySerAlaLeuAspLysI 359 :::	Db 3192 CTTCGGTGCTCTCGCAAGTTTGTTCTCTAATGTTCCTGGTGACGAGCTCGTAAGTCTTGA 3251 Qy 323 323 Db 3252 TCTCTATCCCAATCATCTCTTCCTTATCAATTGAACTCTTTTCTTTAATGCTGGC 3311 Qy 324AlaGlyLysArgIleGlyValPheSerTyrGlySerGlyLeuAlaA 339 Db 3312 TTTCTCTTGAACAGGTCGGCAAGCGCATTGCTCTTACGCCTACGGATCTGGAGCTGCTG 3371	in in in in

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Alignment Scores:
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Query Match:
DB:
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APPLICATION NUMBER: US/08/998,416
FILING DATE: 24-DEC-1997
CLASSIFICATION: 435
PRIOR APPLICATION: 435
PRIOR APPLICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: CH 0016/97
FILING DATE: 31-DEC-1996
ATTORNEY/AGENT INFORMATION:
NAME: Meigs, J. Timothy
REGISTRATION NUMBER: 98,241
REFERENCE/DOCKET NUMBER: PF/5-30306,
TELECOMMUNICATION INFORMATION:
TELEPHONE: 919-541-8587
TELEPHONE: 919-541-8587
INFORMATION FOR SEQ ID NO: 730:
SEQUENCE CHARACTERISTICS:
LENGTH: 635 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
ORGANISM: PAG1476UP
US-08-998-416-730
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                  TTTGACTCTGTGCGTG
                                                        CGGCCCACTGGCGGTGCGGGAACTGTCGCTCTCCTGATCGGTCCAGACGCCCCCATTGTC
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US-09-833-381-1648

US-09-833-381-1648

Sequence 1648, Application US/09833381

Patent No. 6672186

GENERAL INFORMATION:
APPLICANT: Robison, Keith E.
TITLE OF INVENTION: No. 6672186el Nuclei
FILE REFERENCE: 5800-119
CURRENT APPLICATION NUMBER: US/09/833,38
CURRENT FILING DATE: 2001-04-11
PRIOR APPLICATION NUMBER: 09/516,448
PRIOR FILING DATE: 2000-02-29
NUMBER OF SEQ ID NOS: 2050
SOFTWARE: FastSEQ for Windows Version 3
SEQ ID NO 1648
LENGTH: 472
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(472)
OTHER INFORMATION: n = A,T,C or G
US-09-833-381-1648
            RESULT 13
US-09-107-532A-1715
; Sequence 1715, Application U
; Patent No. 6583275
; GENERAL INFORMATION:
} GENERAL INFORMATION:
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Pred. No.:
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Best Local Similarity:
Query Match:
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        APPLICANT: Lynn A DITITLE OF INVENTION:
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Matches:
Conservative:
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77 MetGluArgAsnAsnLeuSerTyrAspCysIleGlyArgLeuGluValGlyThrGluThr 96 ::: :::	Qy 37 GluLysTyrAspGlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMet 56 :::	17 ValGlyIleValAlaLeuGluIleTyrPheProSerGl ::: 19 ATAGGGATTGATCGTCTTTCCTTATTCCTAATTT	est Local Similarity: 24.82% Mismatches: uery Match: 12.54% Indels: B: Gaps: 4 Gaps:	Scores: 5.16e-29 Length: 1 315.00 Matches: 1 milarity: 40.10% Conservative: 6	LOCATION: (B) LOCATION SEQUENCE DESCRIPTION: SEQ IS-09-107-532A-1715	ORGANISM: Enterococc FEATURE: NAME/KEY: misc_feat	MOLECULE TYPE: DNA (ge- HYPOTHETICAL: NO ANTI-SENSE: NO ORIGINAL SOURCE:	LENGTH: 1167 base portion of the control of the con	TELEPHONE: (781)893-5007 TELEFAX: (781)893-8277 INFORMATION FOR SEQ ID NO: 1715: SEQUENCE CHARACTERISTICS:	E: Ariniello, Pamela ISTRATION NUMBER: 40 ERENCE/DOCKET NUMBER	FILING DATE: 14 May 1998 APPLICATION NUMBER: 60/051571 FILING DATE: July 2, 1997 ATTORNEY/AGENT INFORMATION:	APPLICATION NUMBER: US/ FILING DATE: 30-Jun-199 PRIOR APPLICATION DATA: APPLICATION NUMBER: 60/	COMPUTER: PC OPERATING SYSTEM: <unknown application="" ascii="" current="" data:<="" software:="" th=""><th>COUNTRY: USA ZIP: 02354 COMPUTER READABLE FORM: MEDIUM TYPE: CD/ROM</th><th>ADDRESSEE: GENOME STREET: 100 Beave CITY: Waltham STATE: Massachuse</th><th>NUMBER OF SEQUENCES: 73</th></unknown>	COUNTRY: USA ZIP: 02354 COMPUTER READABLE FORM: MEDIUM TYPE: CD/ROM	ADDRESSEE: GENOME STREET: 100 Beave CITY: Waltham STATE: Massachuse	NUMBER OF SEQUENCES: 73
RESULT 14 US-08-956-171E-364 ; Sequence 364, Application US/08956171E ; Patent No. 6593114 ; GENERAL INFORMATION: ; APPLICANT: Charles Kunsch ; Gil H. Choi	82 CAA	331 PheSerTyrGlySerGlyLeuAlaAlaThrLeuTyrSerLeuLysValT	311 LeuAlaSerValLeuAlaGlnTyrSerProGlnGlnLeuAlaGlyLysArgIleGlyVal 	291 AlaSerLeuLeuValSerAsnGlnAsnGlyAsnMetTyrThrSerSerValTyrGlySer ::: :::	Qy 271 ArgAspValGluLysAlaPheMetLysAlaSerSerGluLeuPheSerGlnLysThrLys 290	Qy 251 SerIleTyrSerGlyLeuGluAlaPheGlyAspValLysLeuGluAspThrTyrPheAsp 270 Db 777 777	Qy 231 LysSerLeuAlaArgMetLeuLeuAsnAspPheLeuAsnAspGlnAsnArgAspLysAsn 250 ::::: ::::: :::: :::: ::::::	Qy 211 PheThrLeuAsnAspPheGlyPheMetIlePheHisSerProTyrCysLysLeuValGln 230	Qy 191 TyrSerValTyrCysLysLysIleHisAlaGlnTrpGlnLysGluGlyAsnAspLysAsp 210 ::: Db 649	Qy 171 TyrProIleValAspGlyLysLeuSerIleGlnCysTyrLeuSerAlaLeuAspArgCys 190 ::: :::	Oy 156MetGlnHisAlaTyrAspPheTyrLysProAspMetLeuSerGlu 170 ::: ::::: Db 532 GACAGCGTATTTCTGACAGAAGATATCTATGATTTCTGGCGTCCAGATTATAGCGAA 588	Qy 153 GlyThrHis 155	Qy 145	Qy 132 GlyThrAlaAlaValPheAsnAlaValAsnTrpIleGlu 144 ::: :::::::::::::::::	Qy 112 PheGluGluSerGlyAsnThrAspIleGluGlyIleAspThrThrAsnAlaCysTyrGly 131	

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AGGTATCGATAAAATAAACTTTTÄCGTTCCAAAGTACTATGTAGACATGGCTAULysTyrAspGlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaili.::: ::: :::	TRANDEDNESS: double OPOLOGY: linear CE DESCRIPTION: SEQ ID NO: 364: 8: 1.24e-25	PRIOR APPLICATION DAY APPLICATION NUMI FILING DATE: Jan ATTORNEY/AGENT INFORM REGISTRATION NUM REGISTRATION NUM REFERENCE/DOCKET REFERENCE/DOCKET TELECOMMUNICATION INFORMATION FOR SEQ ID NO SEQUENCE CHARACTERIST LENGTH: 10813 bo TYPE: nucleic ac	ADDING SYLES
Qy 349 AspAlaThrProGlySerAlaLeuAspLysIleThrAlaSerLeuCysAspLeuLysSer 368		189 ArgCysTyrSerValTyrCysLysLysIleHisAlaGlnTrpGlnLysGluGlyAsnAsp :::::::: 4455 CAAAGCTGGAATGAATACGCAAAA	A

204	SerAlaLeuAspArgCysTyrSerValTyrCysLysIleHisAlaGlnTrpGlnLys 	Оу 185
184 762	ProAspMetLeuSerGluTyrProIleValAspGlyLysLeuSerIleGlnCysTyrLeu	Qу 165 Db 706
705	ATTTTAAAACTTAATGATGATGCCGTAGCATATACTGA	64
164		Qy 154
645	::: GTGGTGAGCCTACTCAAGGTGCCGGTGCAGTTGCAATGATGATTTCACATAACCCAAG	ທ
585 153	SerTrpAspGlyLeuArgGly	Oy 147
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465	GĠTÀTTCAACC	Db 433
125	GluGluSerGlyAsnThrAspIleGluGlyIleAspThr	
112	GlyThrGluThrIleIleAspLysSerLysSerValLysThrAsnLeuMetGlnLeuPhe ::: :::	Qy 93 Db 373
372	ACAGAAGAAGATAAAAAGAAT	Db 328
92	euMetGlu	Оу 76
75 327	MetGlyPheCysThrAspArgGluAspIleAsnSerLeuCysMetThrValValGlnAsn :::::	Qy 56 Db 268
267	LeuGluLysTyrAspGlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLys :::	Qy 36 Db 208
35 207	AspValGlyIleValAlaLeuGluIleTyrPheProSerGlnTyrValAspGlnAlaGlu::::::	Oy 16
	-2 (1-478) x US-09-134-001C-1458 (1-1311)	US-10-622-516
	res: 4.01e-25 Length: 1311 284.50 Matches: 107 arity: 38.70% Conservative: 71 milarity: 23.26% Mismatches: 149 11.33% Indels: 33 Gaps:	Alignment Sco Pred. No.: Score: Percent Simil Best Local Si Query Match: DB:
STAPHYLOCOCCUS	NVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO NVENTION: EPIDERMIDIS FOR DIAGNOSTICS AND THERAPEUTICS ENCE: GTC-007 PLICATION NUMBER: US/09/134,001C LING DATE: 1998-08-13 ICATION NUMBER: US 60/064,964 NG DATE: 1997-11-08 ICATION NUMBER: US 60/055,779 NG DATE: 1997-08-14 SEQ ID NOS: 5674 458 311 Staphylococcus epidermidis C-1458	TITLE OF I FILE REFER CURRENT AP CURRENT FI PRIOR APPL PRIOR FILI PRIOR FILI PRIOR FILI PRIOR FILI PRIOR FILI ORGANISM: US-09-134-001
	Lynn Doucette-Stamm et	APPLICANT

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SerIleAspSerLeuPheGluGlyThrTrpTyrLeuValArgValAspGluLysHisArg 421 ::: AATGCAGATCGTGACATTTTCTATTTAAAATCTATTGATAACAATATTCGT 1290	AlaGluAsnMetLysLeuArgGluAspThrHisHisLeuValAsnTyrIleProGlnGly 401 ::: :: :: ::: GAACATTTTTTCAAACGCTTTGACCAATTAGAATTGAATCATGAACTTGAAAAATCA 1239	AspLeuLysSerArgLeuAspSerArgThrGlyValAlaProAspValPhe 381	LysValThrGlnAspAlaThrProGlySerAlaLeuAspLysIleThrAlaSerLeuCys 364	GlyLysArgIleGlyValPheSerTyrGlySerGlyLeuAlaAlaThrLeuTyrSerLeu 344 ::: ::: GGACAAACGATTGGTCTCTTTAGTTATGGTTCTGGTTCTGTAGGCGAGTTCTTTAGT 1089	SerSerValTyrGlySerLeuAlaSerValLeuAlaGlnTyrSerProGlnGlnLeuAla 324 :::	PheSerGlnLysThrLysAlaSerLeuLeuValSerAsnGlnAsnGlyAsnMetTyrThr 304	GluAspThrTyrPheAspArgAspValGluLysAlaPheMetLysAlaSerSerGluLeu 284 :::::	GlnAsnArgAspLysAsnSerIleTyrSerGlyLeuGluAlaPheGlyAspValLysLeu 264	TyrCysLysLeuValGlnLysSerLeuAlaArgMetLeuLeuAsnAspPheLeuAsnAsp 244:::: ::: ::: ::: :::::::::	GluGlyAsnAspLysAspPheThrLeuAsnAspPheGlyPheMetIlePheHisSerPro 224 ::: ::: TATGCACGTCGCCATAATAAAACACTCGCTGATTTCGCTTCACTATGTTTCCATGTACCA 849	AAGTCATTCCAAGAAAGTTGGAATGAA 789
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Search completed: June 24, 2004, 16:00:40 Job time: 178 secs

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RESULT 1

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Aydroxymethylglutaryl-CoA synthase (EC 4.1.3.5), cytosolic, adrenal isoform - human hydroxymethylglutaryl-CoA synthase

C;Species: Homo sapiens (man)

C;Date: 24-Jul-1998 #sequence_revision 24-Jul-1998 #text_change 05-May-2000

C;Accession: \$45497; 139355

R;Rokosz, L.L.; Boulton, D.A.; Butkiewicz, E.A.; Sanyal, G.; Cueto, M.A.; Lachance, P.A.
Arch. Biochem. Biophys. 312, 1-13, 1994

A;Title: Human cytoplasmic 3-hydroxy-3-methylglutaryl coenzyme A synthase: expression, F.
A;Reference number: 139355; MUID:94304197; PMID:7913309

A;Accession: \$45497

A;Molecule type: mRNA
A;Cross-references: EMBL:L25798; NID:9410027; PIDN:AAA62411.1; PID:9410028

A;Experimental source: fetal adrenal

C;Function:
A;Description: catalyzes the condensation of acetyl coenzyme A (Ac-CoA) with acetoacetyl tase

C;Superfamily: hydroxymethylglutaryl-CoA synthase

C;Superfamily: hydroxymethylglutaryl-CoA synthase

C;Keywords: carbon-carbon lyase; cholesterol biosynthesis; coenzyme A; oxo-acid-lyase

F;129/Active site: Cys (covalent substrate-binding) #status predicted S 멍 S 밁 S g B S S S B 8 Query Page 14 Best La Match 98.8%; ocal Similarity 91.9%; s 478; Conservative 301 259 199 181 121 241 61 61 FGDVKLEDTYFDRDVEKAFMKASSELFSQKTKASLLVSNQNGNMYTSSVYGSLASVLAQY EGIDTTNACYGGTAAVFNAVNWIESSSWD----EGIDTTNACYGGTAAVFNAVNWIESSSWDGRYALVVAGDIAVYATGNARPTGGVGAVALL Score 2480; DB 2; Pred. No. 4.4e-177; 0; Mismatches 0; Length 520; Indels 42; Gaps 318 expression, 180 120 258 240 198 120 360 300 149 60 60 1; acetoacetyl P.A.,

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3-hydroxy-3-methy

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RESULT 2
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hydroxymethylglutaryl-CoA synthase (EC 4.1.3.5), cytosolic, fib C; Species: Homo sapiens (man)
C; Date: 07-Apr-1994 #sequence_revision 07-Apr-1994 #text_change C; Accession: $27197; $21590
R; Russ, A.P.; Ruzicka, V.; Maerz, W.; Appelhans, H.; Gross, W. Biochim. Biophys. Acta 1132, 329-331, 1992
A; Title: Amplification and direct sequencing of a cDNA encoding A; Reference number: $27197; MUID:93041939; PMID:1358203
A; Accession: $27197
A; Molecule type: mRNA
A; Residues: 1-520 < RUS>
A; Cross-references: EMBL:X66435; NID:g30008; PIDN:CAA47061.1; FC; Function:
A; Description: catalyzes the condensation of acetyl coenzyme A case
C; Superfamily: hydroxymethylglutaryl-CoA synthase
C; Keywords: carbon-carbon lyase; cholesterol biosynthesis; coer F;129/Active site: Cys (covalent substrate-binding) #status pre
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Similarity 90.9%;
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hydroxymethylglutaryl-CoA synthase (EC 4.1.3.5), cytosolic - rat C; Species: Rattus norvegicus (Norway rat)
C; Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change (C; Accession: S12736
R; Ayte, J.; Gil-Gomez, G.; Hegardt, F.G.
Nucleic Acids Res. 18, 3642, 1990
A; Title: Nucleotide sequence of a rat liver cDNA encoding the cyt A; Reference number: S12736; MUID:90301491; PMID:1972979
A; Accession: S12736
A; Molecule type: mRNA
A; Residues: 1-520 < AYT>
A; Cross-references: EMBL:X52625; NID:g55946; PIDN:CAA36852.1; PID C; Superfamily: hydroxymethylglutaryl-CoA synthase C; Keywords: carbon-carbon lyase; coenzyme A; cytosol; oxo-acid-ly
hydroxymethylglutaryl-CoA synthase (EC 4.1.3.5) - Chinese C; Species: Cricetulus griseus (Chinese hamster) C; Date: 30-Jun-1988 #sequence_revision 30-Jun-1988 #text_C; Accession: A25332 R; Gil, G.; Goldstein, J.L.; Slaughter, C.A.; Brown, M.S. J. Biol. Chem. 261, 3710-3716, 1986 A; Title: Cytoplasmic 3-hydroxy-3-methylglutaryl coenzyme A; Reference number: A25332; MUID:86140166; PMID:2869035 A; Accession: A25332 A; Molecule type: mRNA A; Residues: 1-520 < GIL>
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Symmethylglutaryl-CoA synthase (EC; Species: Gallus gallus (chicken)
C; Species: Gallus gallus (chicken)
C; Date: 04-Dec-1992 #sequence_revisic
C; Date: 04-Dec-1992 #sequence_revisic
C; C; Accession: S13887
R; Kattar-Cooley, P.A.; Wang, H.H.L.;
Arch. Biochem. Biophys. 283, 523-529,
A; Title: Avian liver 3-hydroxy-3-meth
A; Reference number: S13887; MUID: 9111
A; Accession: S13887
A; Molecule type: mRNA
A; Residues: 1-522 < KAT>
A; Note: the authors translated the cc
C; Superfamily: hydroxymethylglutaryl-
C; Keywords: carbon-carbon lyase; coer
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ies: Gallus gallus (chicken)
: 04-Dec-1992 #sequence_revision 04-Dec-1992
ssion: S13887
:ar-Cooley, P.A.; Wang, H.H.L.; Mende-Mueller
Biochem. Biophys. 283, 523-529, 1990
le: Avian liver 3-hydroxy-3-methylglutaryl-Co
srence number: S13887; MUID:91112772; PMID:19
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27-Nov-1997 #sequence revision 12-Dec-1997 #text_c
ion: S71623; A55729; S51103
o, C.; Buesa, C.; Ortiz, J.A.; Haro, D.; Hegardt,
ochem. Biophys. 317, 385-390, 1995
Molecular cloning and tissue expression of human
nce number: S71623; MUID:95200282; PMID:7893153
                                                                                                                                                                 references:
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Addroxymethylglutaryl-CoA synthase (EC 4.1.3.5) precursor - rat
C;Species: Rattus norvegicus (Norway rat)
C;Becies: Rattus norvegicus (Norway rat)
C;Cacession: A35865, 32477
C;Accession: A35865, 32477
R;Ayte, J;Gil-Gomez, G;Haro, D;Marrero, P.F.; Hegardt, P.G.
R;Ayte, J;Gil-Gomez, G;Haro, D;Marrero, P.F.; Hegardt, P.G.
A;Title: Rat mitochondrial and cytosolic 3-hydroxy-3-methylglutaryl-CoA synthases are A;Reference number: A35865;MUID:90251660;PMID:1971108
A;Gitle: Per latiniary
A;Arcserice number: BS:M33648; NID:9204618; PIDN:AAA41336.1; PID:9204619
A;Accession: A35865;MUID:90251660;PMID:1971108
A;Accession: A55865;MUID:90251660;PMID:1971108
A;Accession: C;Ayte, J;Hegardt, F.G.
Bur: J;Hegardt, F.G.
A;Comez, G;Ayte, J;Hegardt, F.G.
R;Gil-Gomez, G;Ayte, J;Hegardt, F.G
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A;Title: Human mitochondrial HMG CoA synthase: liver cDNA and partial A;Reference number: A55729; MUID:95154824; PMID:7851882

A;Accession: B55729

A;Status: preliminary; nucleic acid sequence not shown

A;Molecule type: mRNA

A;Residues: 1-471 <BOU>
A;Cross-references: GB:U12790; GB:U12791

C;Superfamily: hydroxymethylglutaryl-CoA synthase
C;Keywords: carbon-carbon lyase; coenzyme A; mitochondrion: oxo-acid-1
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B55729
hydroxymethylglutaryl-CoA synthase (EC 4.1.3.5), mitochondrial C;Species: Mus musculus (house mouse)
C;Date: 21-Jul-1995 #sequence_revision 21-Jul-1995 #text_change C;Accession: B55729
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erfamily: hydroxymethylglutaryl-CoA synthase
words: carbon-carbon lyase; coenzyme A; mitochondrion;
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Similarity 59.4%; Pred. No. 1.6e-103;
76; Conservative 74; Mismatches 73;
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RESULT 9
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hydroxymethylglutaryl-CoA synthase (EC 4.1.3.5)
C;Species: Blattella germanica (German cockroach
C;Date: 07-Oct-1994 #sequence_revision 01-Dec-19
C;Accession: $38986
R;Martinez-Gonzalez, J.; Buesa, C.; Piulachs, M.
Eur. J. Biochem. 217, 691-699, 1993
A;Title: 3-Hydroxy-3-methylglutaryl-coenzyme-A sa;Reference number: $38986; MUID:94039108; PMID:
A;Rccession: $38986
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-453 <MAR>
A;Cross-references: EMBL:X73679; NID:9416168; PIC;Superfamily: hydroxymethylglutaryl-CoA synthase
C;Keywords: carbon-carbon lyase; coenzyme A; oxc
RESULT 10
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hydroxymethylglutaryl-Configuration  
C; Species: Blattella genometric  
C; Date: 07-Oct-1994 #secondary  
C; Accession: A53565
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RESULT 11
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hydroxymethylglutaryl-Co
N;Alternate names: 3-hyd
C;Species: Pinus sylvest
C;Date: 16-Jul-1999 #seq
C;Accession: T09688
R;Wegener, A.; Gimbel, W
Biochim. Biophys. Acta 1
A;Title: Molecular cloni
A;Reference number: Z168
A;Accession: T09688
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A;Residues: 1-474 <WEG>
A;Cross-references: EMBL
A;Experimental source: t
C;Function:
A;Description: catalyzes
tase
C;Superfamily: hydroxyme
C;Keywords: carbon-carbo
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Molecular cloning of ozone-inducible protein from Pinus sylvertris nce number: Z16823; MUID:97214637; PMID:9061017
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Chem. 269, 11707-11713, 1994
Blattella germanica has two HMG-CoA synthase genes.
Ice number: A53565; MUID:94216267; PMID:7909314
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y: hydroxymethylglutaryl-CoA synthase carbon-carbon lyase; coenzyme A; oxo-

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T49718
probable hydroxymethylglutaryl-CoA synthase
N;Alternate names: protein B23L21.310
C;Species: Neurospora crassa
C;Date: 02-Jun-2000 #sequence_revision 02-J
C;Accession: T49718
R;Schulte, U.; Aign, V.; Hoheisel, J.; Bran
submitted to the Protein Sequence Database,
A;Reference number: Z25022
A;Accession: T49718
A;Recession: T49718
A;Residues: 1-454 <SCH>
A;Cross-references: EMBL:AL356172; GSPDB:GN
A;Experimental source: BAC clone B23L21; st
C;Genetics:
A;Gene: NCSP:B23L21.310
A;Map position: 6
A;Introns: 20/3; 55/1; 409/3
C;Superfamily: hydroxymethylglutaryl-CoA sy
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T09341
hydroxymethylglutaryl-CoA synthase (EC 4.1.3.5) - Arabidopsis thaliana
hydroxymethylglutaryl-coenzyme A synthase; protein T20
C; Species: Arabidopsis thaliana (mouse-ear cress)
C; Species: Arabidopsis thaliana (mouse-ear cress)
C; Pates: 11-Jun-1999 #sequence revision 11-Jun-1999 #text_change 20-Jun-2000
C; Accession: T09341; JC4567
R; Bevan, M.; Hilbert, H.; Braun, M.; Holzer, E.; Brandt, A.; Duesterhoeft, A.;
Submitted to the Protein Sequence Database, June 1999
A; Reference number: Z16650
A; Reperimental source: cultivar Columbia; BAC clone T26M18
A; Residues: 1-461 CABEV
A; Cross-references: EMBL:AL078606
A; Experimental source: cultivar Columbia; BAC clone T26M18
R; Monitamat, F.; Guilloton, M.; Karst, F.; Delrot, S.
Gene 167, 197-201, 1995
Gene 167, 197-201, 1995
A; Title: Isolation and characterization of a cDNA encoding Arabidopsis thalia:
A; Reference number: JC4567; MUID:96144274; PMID:8586777
A; Accession: JC4567
A; Accession: JC4567; MUID:96144274; PMID:8586777
A; Accession: JC4567; MUID:96144274; PMID:8586777
A; Accession: JC4567
A; Cross-references: EMBL:X33882; NID:91143389; PIDN:CAA58763.1; PID:91143390
C; Comment: This enzyme mediates the conversion of three acetyl-CoA molecules to ivity, and uptakes amino acids.
C; Gene: ATSP:T26M18.30
A; Gene: ATSP:T26M18.30
A; Gene: ATSP:T26M18.30
A; Map position: 4
A; Introns: 21/3; 57/2; 101/3; 149/3; 202/3; 227/2; 252/3; 271/2; 300/3; 347/3.
C; Superfamily: hydroxymethylglutaryl-CoA synchase
C; Keywords: carbon-Carbon lyase; coenzyme A; glycoprotein; oxo-acid-lyase
F;117/Accive site: Cys (covalent substrate-binding) #status predicted
F;269/Binding site: carbohydrate (Asn) (covalent) #status predicted
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ALIGNMENTS

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Pfam; PF01154; HMG_COA_synt; 1.
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Hypothetical protein.
SEQUENCE 520 AA; 57569 MW; E28F87
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RESULT 3
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DT 01-OCT-2002 (TrEMBLrel. 22, Created)
DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
GN HMGCS1.
OS Mus musculus (Mouse).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Eut
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mu
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Retina;
RA Strausberg R.;
SLAH31363.1; -.
DR MGD; MGI:107592; Hmgcs1.
DR GO; GO:0004421; F:hydroxymethylglutaryl-CoA synthase ac
DR GO; GO:0004421; F:hydroxymethylglutaryl-CoA synthase ac
DR GO; GO:0006084; P:accetyl-CoA metabolism; IEA.
DR InterPro; IPR000590; HMG_COA_synth.
DR InterPro; IPR000590; HMG_COA_synth.
DR Pfam; PF01154; HMG_COA_SYNTHASE; 1.

Euteleostomi;
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Q1-JUN-2003 (TrEMBLrel. 24, Created)
O1-JUN-2003 (TrEMBLrel. 24, Last sequence update)
O1-OCT-2003 (TrEMBLrel. 25, Last annotation update)
Hypothetical protein.

Enactiopheral protein.
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Caryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleosto
Cinopterygii; Neopterygii; Teleostei; Ostariophysi; Cyprini
Crinidae; Danio.
                          al Similarity
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APESVLASTANEHFPSPAKKVPRIP-PAAEAEPISVTNGEH 520
WPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCTDREDINSLCMTV
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CSLEDFGFMVFHSPYCKLVQKSLARLMLNDFLCHPSPNMESGPFSGLEAFRDVKIEDTYF
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   RP SEQUENCE FROM N.A.

STRAIN-C57BL/6J; TISSUE-Liver;

RC STRAIN-C57BL/6J; TISSUE-Liver;

RA MEDLINE-21085660; PubMed=11217851;

RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,

RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,

RA Arakawa T., Hara A., Fukunishi Y., Konno H., Kadchi J., Fukuda S.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kasukawa T., Saito R.,

RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kasukawa T., Saito R.,

RA Aizawa K., Izawa M., Nishi K., Nisaido I., Pesole G., Quackenbush J.,

Ra Kadota K., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,

Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,

RA Sakai H., Staubli F., Suzuki R., Carninci P., de Bonaldo M.F.,

RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,

ROSTINIC P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,

RA Mordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,

Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,

RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,

"Functional annotation of a full-length mouse cDNA collection.";

Nature 409:685-690(2001).

PSEQUENCE FROM N.A.
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Q9DBM4;
Q9DBM4;
Q1-JUN-2001 (TrEMBLrel. 17, Created)
01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
3-hydroxy-3-methylglutaryl-coenzyme A synthase 2.
                                                                                                                                                                                                                                                                                  Mus musculus (Mouse).
Eukaryota; Metazoa; Chordata;
Mammalia; Eutheria; Rodentia;
NCBI_TaxID=10090;
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; Murinae; Mus.
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Strausberg R.;

Submitted (MAR-2002) to the EMBL/GenBank/DI

EMBL; AK004865; BAB23626.1; -.

EMBL; BC014714; AAH14714.1; -.

EMBL; BC024744; AAH24744.1; -.

PIR; B55729; B55729.

MGD; MGI:101939; Hmgcs2.

GO; GO:0004421; F:hydroxymethylglutaryl-Col

GO; GO:0004421; F:hydroxymethylglutaryl-Col

GO; GO:0006084; P:acetyl-CoA metabolism; II

InterPro; IPR000590; HMG_CoA_synth.

InterPro; IPR000590; HMG_CoA_synt; I.

Pfam; PF01154; HMG_COA_synt; 1.

PROSITE; PS01226; HMG_COA_SYNTHASE; 1.

SEQUENCE 508 AA; 56822 MW; 5A20ACE918FI
Q9DBK1;
Q9DBK1;
Q9DBK1;
Q1-JUN-2001 (TrEMBLre)
Q1-JUN-2001 (TrEMBLre)
Q1-OCT-2003 (TrEMBLre)
Mus musculus (Mouse).
Eukaryota; Metazoa; C)
Mammalia; Eutheria; Ro
NCBI_TaxID=10090;
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STRAIN=C57BL/6J; TISSIMEDLINE=21085660; Publication of the company of the comp
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Putative CG43
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Kadota K., Matsuda H.A., Ashburner M., Kasukawa T., Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casav Fleischmann W., Gaasterland T., Gissi C., King B., Kochi Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Qu Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner I Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., E Blake J., Boffelli D., Bojunga N., Carninci P., de Bonal Brownstein M.J., Bult C., Fletcher C., Fujita M., Garibo Gustincich S., Hill D., Hofmann M., Hume D.A., Kamiya M. Lyons P., Marchionni L., Mashima J., Mazzarelli J., Momk Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamotc Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Suzuki H., Toyo-oka K., Wang K.H., Weitz C., Whittaker C Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Ko Hayashizaki Y., Mang K.H., Weitz C., Whittaker C Mature 409:685-690(2001).

RMBL; AKO04902; BAB23657.1; -.
RMGD; GO:0006084; P:acetyl-CoA metabolism; IEA.
InterPro; IPR008260; HMG_COA_synth.
InterPro; IPR008260; HMG_COA_synth.
RPGOSITE; PS01226; HMG_COA_synthase; 1.
Pfam; PF01154; HMG_COA_synt; 1.
PS01126; HMG_COA_SYNTHASE; 1.
SEQUENCE 508 AA; 56822 Mw; 0F20AF327392F183 CRC64;
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"Finctional T., Washida K., Hasegawa Y., Kawaji H., Kohtsuki S., Hayashizaki Y.;
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3.2e-101;
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2000 (TrEMBLrel. 13, Created)
2000 (TrEMBLrel. 13, Last sequence update)
2003 (TrEMBLrel. 25, Last annotation update)
2 CG4311 protein (CG16796 protein) (LD26976P)
CG4311 OR CG16796.

PRELIMINARY;

PRT;

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                                                                                        Denocophila melanogastor (Fruit fly)

Chepters, Endopteryora, Metazol, atthropoda, Haxapoda, Insecta, Ptaygota,

Chepters, Endopteryora, Dipters, Brachycera, Muscomorpha,

Chepters, Brachycer, Brachycer, Chepters, Brachycer, Chepters, M. Hookin, M.
                                 Query Match
Best Local Si
Matches 263;
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WPKDVGIVALEIYFPSQYVDQAELEKYDGVDAGKYTIGLGQAKMGFCTDREDINSLCMTV
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larity 57.3%;
Conservative
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                             ore 1337; DB seed. No. 6e-90; Mismatches
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Q9NDA8;
01-OCT-2000 (TrEMBLrel. 15, C)
01-OCT-2000 (TrEMBLrel. 15, I)
01-OCT-2003 (TrEMBLrel. 25, I)
3-hydroxy-3-methylglutaryl cc
                                                                                                                                                                   Dendroctonus jeffreyi (Jeffrey pine beetle).
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygot.
Neoptera; Endopterygota; Coleoptera; Polyphaga; Cucujiformi.
Phytophaga; Scolytidae; Dendroctonus.
NCBI_TaxID=77162;
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                 Local Similarity 50.4%; les 231; Conservative 6
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Pred. No. 3.2e-77;
64; Mismatches 109;
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Q8H051;
01-MAR-2003 (TrEMBLrel. 23, Created)
01-MAR-2003 (TrEMBLrel. 23, Last sequence up
01-OCT-2003 (TrEMBLrel. 25, Last sequence up
01-OCT-2003 (TrEMBLrel. 25, Last annotation
Putative hydroxymethylglutaryl coenzyme A sy
OJ1263H11.4.

Oryza sativa (japonica cultivar-group).
Eukaryota; Viridiplantae; Streptophyta; Embi
Spermatophyta; Magnoliophyta; Liliopsida; Po
Ehrhartoideae; Oryzeae; Oryza.

NCBI_TaxID=39947;
[1]
SEQUENCE FROM N.A.
Wing R.A., Yu Y., Soderlund C., Kim H.-R., F
Collura K., McCombie D.W.R., de la Bastide N
Kirchoff K., Kuit K., Nascimento L., Zutavel
Baker J., Santos L., Miller B., Katzenbergel
Yang C., O'Shaugnessy A., Palmer L., Dedhia
"Rice Genomic Sequence.";
                                                                                                                                                                                                   "Rice Genomic Sequence.";
Submitted (JAN-2003) to the EMBL/GenBank/DDBJ data
EMBL; AC118980; AAO15287.1; -.
GO; GO:0004421; F:hydroxymethylglutaryl-CoA syntha
GO; GO:0006084; P:acetyl-CoA metabolism; IEA.
InterPro; IPR0008260; HMG_CoA_synth.
InterPro; IPR000590; HMG_CoA_synt_AS.
Pfam; PF01154; HMG_COA_synt; 1.
PROSITE; PS01226; HMG_COA_SYNTHASE; 1.
SEQUENCE 463 AA; 51502 MW; 60D021C5C48E99B6 CF
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Pred. No. 1.3e
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la Bastide M., Spiegel L., Preston
o L., Zutavern T., Balija V., Bell I
Katzenberger F., Muller S., King L
r L., Dedhia N.;
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Q1-MAR-2001 (TrEMBLrel. 16, Created)
C1-MAR-2001 (TrEMBLrel. 16, Last sequence update)
C1-OCT-2003 (TrEMBLrel. 25, Last annotation update)
Putative 3-hydroxy-3-methylglutaryl coenzyme A synthase.
Phycomyces blakesleeanus.
Eukaryota; Fungi; Zygomycota; Zygomycetes; Mucorales; Muc
                                                                                                                                                                                                                                                                                              ry Match 41.6%;
t Local Similarity 47.5%;
ches 217; Conservative (
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3-hydroxy-3-methylglutaryl
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Q8C5F4 PRELIMINARY; PRT; 268 AA.
Q8C5F4;
01-MAR-2003 (TrEMBLrel. 23, Created)
01-MAR-2003 (TrEMBLrel. 23, Last sequence updated)
01-OCT-2003 (TrEMBLrel. 25, Last annotation upout transcription factor 1.
HMGCS1 OR B130032C06RIK.
Mus musculus (Mouse).
Eukaryota; Metazoa; Chordata; Craniata; Verteb:
Mammalia; Eutheria; Rodentia; Sciurognathi; Mu:
NCBI_TaxID=10090;
[1]
SEQUENCE FROM N.A.
STRAIN=C57BL/6J; TISSUE=Testis;
MEDLINE=22354683; PubMed=12466851;
The FANTOM Consortium,
the RIKEN Genome Exploration Research Group Ph.
"Analysis of the mouse transcriptome based on 60,770 full-length cDNAs.";
Nature 420:563-573 (2002).
EMBL; AK078743; BAC37373.1; -.
MGD; MGI:107592; Hmgcsl.
G0; G0:0004421; F:hydroxymethylglutaryl-CoA sy.
G0; G0:0004421; F:hydroxymethylglutaryl-CoA sy.
G0; G0:0004421; F:hydroxymethylglutaryl-CoA sy.
G0; G0:0006084; P:acetyl-CoA metabolism; IEA.
InterPro; IPR000590; HMG_COA_synth.
InterPro; IPR000590; HMG_COA_synth.
InterPro; IPR000590; HMG_COA_synth.
PROSITE; PS01226; HMG COA_SYNTHASE; 1.
PROSITE; PS01226; HMG COA_SYNTHASE; 1.
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C STRAIN=cv. RRIM600;

R Hallahan D.L., Keiper-Hrynko N.M.;

Hallahan D.L., Keiper-Hrynko N.M.;

Hallahan D.L., Keiper-Hrynko N.M.;

R Genes involved in the biosynthesis of isopentenyl diphosphate in rubber tree Hevea brasiliensis.";

Submitted (OCT-2001) to the EMBL/GenBank/DDBJ databases.

EMBL; AF429389; AAL18930.1; -.

R GO; GO:0004421; F:hydroxymethylglutaryl-CoA synthase activity; IEAR

GO; GO:0004421; F:hydroxymethylglutaryl-CoA synthase activity; IEAR

GO; GO:0004421; F:hydroxymethylglutaryl-CoA synthase activity; IEAR

GO; GO:0004421; F:hydroxymethylglutaryl-CoA synthase activity; IEAR

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01-DEC-2001 (TrEMBLrel. 19, Last sequence update)
01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
11-OCT-2003 (TrEMBLrel. 25, Last annotation update)
Hydroxymethylglutaryl coenzyme A synthase.
Hevea brasiliensis (Para rubber tree).
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosid;
eurosids I; Malpighiales; Euphorbiaceae; Crotonoideae; Micrandreae
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ALIGNMENTS

ACCESSION VERSION KEYWORDS SOURCE RESULT 1 AK045094 LOCUS DEFINITION MS 2 Mus musculus Eukaryota; Me Mammalia; Eut Mus musculus 9.5 days embryo parthenogenote cDNA, RIKEN full-length enriched library, clone:B130032C06 product:pre B-cell leukemia transcription factor 1, full insert sequence.

AK045094
AK045094.1 GI:26337068
HTC; CAP trapper.
Mus musculus (house mouse) ; Metazoa; Chordata; Butheria; Rodentia; Craniata; Vertebrata; E Sciurognathi; Muridae; Euteleostomi;
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PANTON Consortium and the RIKEN Genome Exploration Research Nature 409, 685-690 (2001)

E The PANTON Consortium and the RIKEN Genome Exploration Research Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs

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High-efficiency full-length cDNA cloning

L Meth. Enzymol. 303, 19-44 (1999)

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Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.

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hai,T., Tashiro,H., Itoh,M.,
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shizaki, The Institute of, Laboratory for Genome mic Sciences Center (GSC), ro-cho, Tsurumi-ku, Yokohama, e-res@gsc.riken.go.jp, l:81-45-503-9222,

uArg	Qy 41 GlyValAspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMetGlyPheCysThr 60	QY 21 AlaLeuGluIleTyrPheProSerGlnTyrValAspGlnAlaGluLeuGluLysTyrAsp 40 	Qy 1 MetProGlySerLeuProLeuAsnAlaGluAlaCysTrpProLysAspValGlyIleVal 20	: 11 Gaps: 1 -10-622-516-2 (1-478) x AK044835 (1-3286)	t s oca	GIN gnment Scores: d. No.:	<pre>polyA_signal 32723277 /note="putative" polyA_site 3286 /note="putative"</pre>		SVKSNLMQLFEESGNTDIEGIDTTNACYGGTAAVFNAVNWVESSSWDGRYALVVAGDI AIYATGNAR PTGGVGAVALLIGPNAPLIFDRGLRGTHMQHAYDFYKPDMLSEYPVVDG KLSIQCYLSALDRCYSVYRKKIRAQWQKEGKDKDFTLNDFGFMIFHSPYCKLVQKSLA RMFLNDFLNDONRDKNSIYSGLFAFGDVKLEDTVEDDDVEVAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGA	/ COUNT_BCAIL=1 / protein_id="BAC32112.1" / db_xref="GI:26336857" / translation="MPGSLPLNAEACWPKDVGIVALEIYFPSQYVDQAELEKYDGVDA GKYTIGLGOARMGFCTDAEDTNGIGIGIGIGIGIGIGIGIGIGIGIGIGIGIGIGIGIGI	transcription factor 1 (MGD MGI:97495, GB NM_008783, evidence: BLASTN, 99%, match=124) putative"	one_lib="RIKEN full-length enriched mous v_stage="9.5 days embryo" 1685	xref="MGI:2410190" xref="taxon:10090" one="B130007E24"	n="Mus muscul)="mRNA" 'C57BL/6J" 'FANTOM DB:B	ip://genome.g ip://fantom.g Location/Q	ience, Tokyo University of Agriculture, 1 nagawa Prefecture, Japan) whose assistanc knowledge.	enomic Sciences Center and Genome Exploration Research Grouivision of Experimental Animal Research in Riken contrirepare mouse tissues.	ax:81-45-503-9216) DNA library was prepared and sequenced in Mouse Genome
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Please visit our web site for further details.
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Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.go.jp, URL:http://genome.gsc.riken.go.jp/, Tel:81-45-503-9222, Fax:81-45-503-9216)
cDNA library was prepared and sequenced in Mouse Genome
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Alignment Scores: Pred. No.: 1.36e-162 Length: Score: 1513.00 Matches: 282	source 11527 /organism="Homo sapiens" /mol_type="genomic DNA" /db_xref="taxon:9606" spene	Adams, M.D. and Cargill, M. Direct Submission Submitted (16-NOV-2003) Celera Genomics, 45 W. Rockville, MD 20850, USA This sequence was made by sequencing genomic of them based on alignment. Location/Qualifiers	g nonneutral os 302 (5652), 8 1 to 1527) G., Glanowsk ., Tanenbaum	Eukaryota; Metazoa; Chordata; Craniata; Ve Eukaryota; Metazoa; Chordata; Craniata; Ve Mammalia; Eutheria; Primates; Catarrhini; 1 (bases 1 to 1527) 5 Clark, A.G., Glanowski, S., Nielson, R., Thom Todd, M.A., Tanenbaum, D.M., Civello, D.R., L Ferriera, S., Wang, G., Zheng, X.H., White, T. Adams, M.D. and Cargill, M.	6670 sapien mic sur 6670 6670.1		399 ProGlnGlySerIleAspSerLeuPheGluGlyThrTrpTyrLeuValArgValAspGlu	339 AlaThrLeuTyrSerLeuLysValThrGlnAspAlaThrProGlySerAlaLeuAspLys

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Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A., Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B., Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J., Ferriera, M.D. and Cargill, M.
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This sequence was made by sequencing
them based on alignment.
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Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A., Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B., Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J., Adams, M.D. and Cargill, M.
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Group Phase I & II Team.

Analysis of the mouse transcriptome based on functional annotation of 60,770 fulll-length cDNAs

Analysis of the mouse transcriptome based on functional annotation of 60,770 fulll-length cDNAs

All Nature 420, 563-573 (2002)

CE 6 (bases 1 to 3287)

RS Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H., Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y.,

Furuno, M., Hanagaki, T., Hara, A., Hayarsu, M., Hiramoto, K.,

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Kasukawa, T., Kato, H., Kawai, J., Kojima, Y., Konno, H., Koda, M.,

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Nomura, K., Numazaki, R., Ohno, M., Okadai, Y., Koido, T., Owa, C.,

Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D.,

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NSLCLTVVQRLMERTKLPWDAVGRLEVGTETIIDKSKAVKTVLMELFQDSGNTDIEGI
DTTNACYGGTASLFNAANWMESSYWDGRYALVVCGDIAVYPSGNARPTGGAGAVAMLI
GPKAPLVLEQGLRGTHMENAYDFYKPNLASEYPLVDGKLSIQCYLRALDRCYAAYRKK
IQNQWKQAGNNQPFTLDDVQYMIFHTPFCKMVQKSLARLMFNDFLSSSSDKQNNLYKG
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                           Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y. Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new Genome Res. 10 (10), 1617-1630 (2000)

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Functional annotation of a full-length mouse Nature 409, 685-690 (2001)
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                                                           /organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="FANTOM_DB:13000(
/db_xref="MGI:1910606"
/db_xref="taxon:10090"
/clone="1300004J23"
/sex="male"
  /note="unnamed protein product;
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                           tissue_type="liver"
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Full-length cDNA libraries and nor
Unpublished (2001)
Contact: Genoscope
Genoscope - Centre National de Seq
BP 191 91006 EVRY cedex - France
Email: segref@genoscope.cns.fr, We
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/mol_type="mRNA"
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/clone="CSODF016YJ04"
/tissue_type="FETAL BRAIN"
/dev_stage="fetal"
/clone_lib="Homo sapiens FETAL BRAIN"
/note="Organ: brain; Vector: pCMVSPORT_6; 1st strand cDNA
was primed with a NotI-oligo(dT) primer. Five prime end
enriched, double-strand cDNA was digested with Not I and
cloned into the Not I and EcoRV sites of the pCMVSPORT 6
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SM Pan troglodytes

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleoston

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.

E 1 (bases 1 to 1420)

S Clark,A.G., Glanowski,S., Nielson,R., Thomas,P., Kejariwal,A.,

Todd,M.A., Tanenbaum,D.M., Civello,D.R., Lu,F., Murphy,B.,

Ferriera,S., Wang,G., Zheng,X.H., White,T.J., Sninsky,J.J.,

Adams,M.D. and Cargill,M.

Inferring nonneutral evolution from human-chimp-mouse orthologous gene trios

Science 302 (5652), 1960-1963 (2003)

D 14671302

E 2 (bases 1 to 1420)

S Clark,A.G., Glanowski,S., Nielson,R., Thomas,P., Kejariwal,A.,

Todd,M.A., Tanenbaum,D.M., Civello,D.R., Lu,F., Murphy,B.,

Ferriera,S., Wang,G., Zheng,X.H., White,T.J., Sninsky,J.J.,

Adams,M.D. and Cargill,M.

Direct Submission

L Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,

Rockville, MD 20850, USA

This sequence was made by sequencing genomic exons and ordering them based on alignment.

Location/Qualifiers

1. 1420

Commission 11420

Commission 1
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Tissue Procurement: ATCC
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
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Location/Qualifiers
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National Institutes of Health, Mammalian
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Contact: Robert Strausberg, Ph.D.
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1 (bases 1 to 1201)

Li,W.B., Gruber,C., Jessee,J. and Polayes,D.

Full-length cDNA libraries and normalization

Unpublished (2001)

Contact: Genoscope

Genoscope - Centre National de Sequencage

BP 191 91006 EVRY cedex - France

Email: segref@genoscope.cns.fr, Web: www.genoscope.cns.fr

Library was constructed by Life Technologies, a division of
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Invitrogen. This sequence belongs to sequence cluster 309 more information about this cluster, see http://www.genoscope.cns.fr/cgi-bin/cluster.cgi?seq=CSODF022BC09QP1&cluster=3098.r. (Feng Liang Email : fliang@lifetech.com URL : http://fulllength.invitrogen.com/ InVitroGen Corporation http://fulllength.invitrogen.com/ InVitroGen Corporation http://sequence Genoscope sequence ID : CSODF022BC09OP1. 3098 Contact 1600 H For

150
570 GCTGTATATGCCACAGGAAATGCTAGACCTACAGGTGGAGTTGGAGCAGTAGCTCTGCTA 629
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141 AsnTrpIleGluSerSerSerTrpAsp 149
121 GluGlyIleAspThrThrAsnAlaCysTyrGlyGlyThrAlaAlaValPheAsnAlaVal 140
101 SerLysSerValLysThrAsnLeuMetGlnLeuPheGluGluSerGlyAsnThrAspIle 120
81 AsnLeuSerTyrAspCysIleGlyArgLeuGluValGlyThrGluThrIleIleAspLys 100
61 AspArgGluAspIleAsnSerLeuCysMetThrValValGlnAsnLeuMetGluArgAsn 80
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1 MetProGlySerLeuProLeuAsnAlaGluAlaCysTrpProLysAspValGlyIleVal 20
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201 nism="Homo cype="mRNA"
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AUTHORS
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                                                                                                                                                               Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLN
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM14058 row: j column: 02
High quality sequence stop: 717.
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National Institutes of Health, Ma
Unpublished (1999)
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/mol_type="mRNA"
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Tissue Procurement: Dr. James Lin,
cDNA Library preparation: Dr. M. B.
CDNA Library Arrayed by: Dr. M. B.
CDNA Sequencing by: Dr. M. Bento S.
Clone Distribution: Researchers management (www.resgen.com).
Seq primer: M13 REVERSE.
Location/Qualifiers
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Rattus.
1 (bases 1 to 789)
1 (bases 1 to 789)
Bonaldo, M.F., Lennon, G.
Normalization and subtra
discovery
Genome Res. 6 (9), 791-8
97044477
8889548
Contact: Soares, MB
Coordinated Laboratory 1
University of Iowa
375 Newton Road, 4156
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                 GlnLysSerLeuAlaArgMetLeuLeuAsnAspPheLeuAsnAspGlnAsnArgAspLys
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/db_xref="taxon:10116"
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/lab_host="DH10B (Life Technologies) (T1 phage resistant)"
/clone_lib="UI-R-FJ0"
/clone_lib="UI-R-FJ0"
/note="Vector: pYX-Asc; Site_1: EcoR I; Site_2: Not I;
/clone_lib="UI-R-FJ0"
/note="Vector: pYX-Asc; Site_1: EcoR I; Site_2: Not I;
/UI-R-FJ0 is a cDNA library containing the following
tissue(s): rat embryo. The library was constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT7T3-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tag for this library is CATCTCTACT. This library
was created for the University of Iowa Program for Rat
Gene Discovery and Mapping (Val Sheffield, Bento Soares
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Bento Soares, University of
Soares, University of Iowa
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E 1 (bases 1 to 779)

R NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov

Tissue Procurement: Dr. Jim Lin, University of Iowa
cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
Clone Distribution: Distribution information can be found at

http://genome.uiowa.edu/distribution/mousefl.html
This clone was contributed by the Brain Molecular Anatomy Project
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Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Susan L. Sullivan, PhD.
cDNA Library Preparation: ResGen, Invitrogen Corp
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM13881 row: n column: 05
High quality sequence stop: 599.
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/mol_type="mRNA"
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/clone="IMAGE:6392260"
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Primer: Oligo dT. Average insert size 2.2 kb. Constructed by ResGen, Invitrogen Corp. Note: this is a NIH_MGC
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920 GGACGNNTAATTTAGAAGATACTTACTTTTGACCGAGATGTAAAAAAG 967	260 yAsp-ValLysLeuGluAspThrTyr-PheAspArgAspValGluLys 275	861 TINTCCTATGATCAAAACAGAGATAA-AACAGTATTTACAGTGGACTGGAAGCCTTTTGG 919	241 PheLeuAsnAspGlnAsnArgAspLysAsnSerIleTyrSerGlyLeuGluAla-PheGl 260	801 TTTCACTCACCATATTTGTAACTGGTGCAGAAATCTCTAGCTCGGATGTCCCTGAATGAC 860	221 PheHisSerProTyrCysLysLeuValGlnLysSerLeuAlaArgMetLeuLeuAsnAsp 240	742 CAGTGGCAGAAAGAGGGAAAGGATAAAGA-TTTACCCTGAATGATNTTGGCTTCATGATC 800	201 GlnTrpGlnLysGluGlyAsnAspLysAspPheThrLeuAsnAspPheGlyPheMetIle 220	682 CAGTGCTACCTCAGCGCCCTGGACCGCTGCTATTCTGTCTACCGCAAAAAAGATCCGTGCC 741	181 GlnCysTyrLeuSerAlaLeuAspArgCysTyrSerValTyrCysLysLysIleHisAla 200	622 GACTTTACAAGCCTGACATGCTCTCCGAGTACCCTGTGGTCGACGGAAAGCTCTCCATA 681	ProlleValAspGlyLysLeuSerI	562 CCAAACGCTCCTAATTTTTGACCGANGGCTCCGTGNGACACACATGCAGCATGCCTAT 621	151LeuArgGlyThrHisMetGlnHisAlaTyr 160	502 TATGCCACAGGAAATGCCAGACCTACAGGTGGAGTTGGAGCTGTGGCCCTGCTAATTGGG 561	150 150	442 GTCGAATCCAGCTCTTGGGATGGACGATATGCTCTGGTAGTTGCAGGAGACATTGCTATA 501	143 IleGluSerSerSerTrpAspGly	382 ATAGATACAACCAATGCATGCTATGGGGGCACAGCTGCAGTCTTCAATGCCGTGAACTGG 441	123 IleAspThrThrAsnAlaCysTyrGlyGlyThrAlaAlaValPheAsnAlaValAsnTrp 142	322 TCAGTGAAGTCTAATTTGATGCAGCTGTTTGAGGAGTCTGGGAATACAGATATAGAAGGA 381	103 SerValLysThrAsnLeuMetGlnLeuPheGluGluSerGlyAsnThrAspIleGluGly 122	262 TCCTATGATTGCATTGGGCGGCTAGAAGTTGGAACAGAGACAATCATCGACAAATCGAAA 321	83 SerTyrAspCysIleGlyArgLeuGluValGlyThrGluThrIleIleAspLysSerLys 102	202 GAAGACATCAACTCTTTTGCCTGACTGTGGTTCAGAAACTGATGGAGAGACATAGCCTT 261	63 GluAspIleAsnSerLeuCysMetThrValValGlnAsnLeuMetGluArgAsnAsnLeu 82	142 GATGCTGGAAAGTATACCATCGGCCTGGGCCAGGCCAGG	43 AspAlaGlyLysTyrThrIleGlyLeuGlyGlnAlaLysMetGlyPheCysThrAspArg 62	82 GAAATCTACTTTCCTTCAATATGTCGATCAAGCTGAGTTGGAAAAATACGATGGTGTA 141

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